

1. Introduction

This report presents an inventory of U.S. anthropogenic greenhouse gas emissions and sinks for the years 1990 through 2022 compiled by the United States government. A summary of source and sink category estimates is provided in Table 2-1, Table 2-2, and Table 2-4 by gas and IPCC sector in the Trends in Greenhouse Gas Emissions and Sinks chapter. The emission and sink estimates in these tables are presented throughout the main report in both CO₂ equivalents (CO₂ Eq.³⁰ and unweighted units). This report also discusses the methods and data used to calculate the emission and sink estimates.

The United States is party to both the 1992 UNFCCC and the 2015 Paris Agreement. The Paris Agreement set a global temperature goal—holding the increase in the global average temperature to well below 2°C above pre-industrial levels and pursuing efforts to limit the increase to 1.5°C—that articulates with greater precision States’ views on what is necessary to meet the UNFCCC’s objective of “stabiliz[ing] ... greenhouse gas concentrations in the atmosphere at a level that would prevent dangerous anthropogenic interference with the climate system.”³¹

The United States is committed to submitting a national inventory of anthropogenic emissions sources and removals by sinks of greenhouse gases by April 15 of each year. The United States has prepared this report, in conjunction with Common Reporting Tables (CRTs) that accompany this report, consistent with its obligations under those agreements.

In 1988, preceding the creation of the UNFCCC, the World Meteorological Organization (WMO) and the United Nations Environment Programme (UNEP) jointly established the Intergovernmental Panel on Climate Change (IPCC). The role of the IPCC is to assess on a comprehensive, objective, open and transparent basis the scientific, technical and socio-economic information relevant to understanding the scientific basis of risk of human-induced climate change, its potential impacts and options for adaptation and mitigation (IPCC 2021). The Paris Agreement and the UNFCCC require use of methods from the *2006 IPCC Guidelines for National Greenhouse Gas Inventories* and encourages Parties to use the *2013 Supplement to the 2006 IPCC Guidelines for National Greenhouse Gas Inventories: Wetlands* and the *2019 Refinement to the 2006 IPCC Guidelines for National Greenhouse Gas Inventories*. The recently released *2019 Refinement* clarify and elaborate on the existing guidance in the *2006 IPCC Guidelines*, along with providing updates to default values of emission factors and other parameters based on updated science. This report applies both the *2013 Supplement* and updated guidance in the *2019 Refinement* to improve accuracy and completeness of the *Inventory*. For more information on specific uses, see Section 1.4 of this chapter on Methodology and Data Sources.

³⁰ More information is provided in the Global Warming Potentials section of this chapter on the use of IPCC *Fifth Assessment Report* (AR5) GWP values.

³¹ See Paris Agreement, Article 2.1(a); UNFCCC, Article 2.

Box 1-1: Methodological Approach for Estimating and Reporting U.S. Emissions and Removals, including Relationship to EPA’s Greenhouse Gas Reporting Program

Consistent with Article 13.7(a) of the Paris Agreement and Article 4.1(a) of the UNFCCC, as well as relevant decisions under those agreements, the emissions and removals presented in this report and this chapter are organized by source and sink categories and calculated using internationally accepted methods in the *2006 IPCC Guidelines for National Greenhouse Gas Inventories (2006 IPCC Guidelines)* and, where appropriate, its supplements and refinements. Additionally, the calculated emissions and removals in a given year for the United States are presented in a common manner in line with the reporting guidelines for the reporting of inventories under the Paris Agreement and the UNFCCC. The Parties’ use of consistent methods to calculate emissions and removals for their inventories helps to ensure that these reports are comparable. The presentation of emissions and removals provided in this *Inventory* does not preclude alternative examinations (e.g., economic sectors). Rather, this *Inventory* presents emissions and removals in a common format consistent with how Parties are to report their national inventories under the Paris Agreement and the UNFCCC. The report itself, and this chapter, follows this common format, and provides an explanation of the application of methods used to calculate emissions and removals.

EPA also collects greenhouse gas emissions data from individual facilities and suppliers of certain fossil fuels and industrial gases through its Greenhouse Gas Reporting Program (GHGRP), which is complementary to the U.S. *Inventory*.³² The GHGRP applies to direct greenhouse gas emitters, fossil fuel suppliers, industrial greenhouse gas suppliers, and facilities that inject carbon dioxide (CO₂) underground for sequestration or other reasons and requires reporting by over 8,000 sources or suppliers in 41 industrial categories.³³ Annual reporting is at the facility level, except for certain suppliers of fossil fuels and industrial greenhouse gases. In general, the threshold for reporting is 25,000 metric tons or more of CO₂ Eq. per year. Facilities in most source categories³⁴ subject to GHGRP began reporting for the 2010 reporting year while additional types of industrial operations began reporting for the 2011 reporting year. Methodologies used in EPA’s GHGRP are consistent with the *2006 IPCC Guidelines*. While the GHGRP does not provide full coverage of total annual U.S. greenhouse gas emissions and removals (e.g., the GHGRP excludes emissions from the Agriculture and Land Use, Land-Use Change, and Forestry sectors), it is an important input to the calculations of national-level emissions in this *Inventory*.

Data presented in this *Inventory* report and EPA’s GHGRP are complementary. The GHGRP dataset continues to be an important resource for the *Inventory*, providing not only annual emissions information, but also other annual information such as activity data and emission factors that can improve and refine national emission estimates and trends over time. Methodologies used in EPA’s GHGRP are consistent with the *2006 IPCC Guidelines* (e.g., higher tier methods). GHGRP data also allow EPA to disaggregate national inventory estimates in new ways that can highlight differences across regions and sub-categories of emissions, along with enhancing the application of QA/QC procedures and assessment of uncertainties. EPA uses annual GHGRP data in several categories to improve the national estimates presented in this *Inventory*, consistent with IPCC methodological guidance. See Annex 9 for more information on specific uses of GHGRP data in the *Inventory* (e.g., natural gas systems).

1.1 Background Information

Science

For over the past 200 years, the burning of fossil fuels such as coal, oil, and natural gas, along with deforestation, land-use changes, and other activities have caused the concentrations of heat-trapping "greenhouse gases" to increase significantly in our atmosphere (IPCC 2021). These gases in the atmosphere absorb some of the energy

being radiated from the surface of the Earth that would otherwise be lost to space, essentially acting like a blanket that makes the Earth's surface warmer than it would be otherwise.

Greenhouse gases are necessary to life as we know it. Without greenhouse gases to create the natural heat-trapping properties of the atmosphere, the planet's surface would be about 60 degrees Fahrenheit cooler than present (USGCRP 2017). Carbon dioxide is also necessary for plant growth. With emissions from biological and geological sources, there is a natural level of greenhouse gases that is maintained in the atmosphere. Human emissions of greenhouse gases and subsequent changes in atmospheric concentrations alter the balance of energy transfers between space and the earth system (IPCC 2021). A gauge of these changes is called radiative forcing, which is a measure of a substance's total net effect on the global energy balance for which a positive number represents a warming effect, and a negative number represents a cooling effect (IPCC 2021). IPCC concluded in its most recent scientific assessment report that it is "unequivocal that human influence has warmed the atmosphere, ocean and land" (IPCC 2021).

As concentrations of greenhouse gases continue to increase from man-made sources, the Earth's temperature is climbing above past levels. The Earth's average land and ocean surface temperature has increased by about 2.0 degrees Fahrenheit from the 1850 to 1900 period to the decade of 2011 to 2020 (IPCC 2021). The last four decades have each been the warmest decade successively at the Earth's surface since at least 1850 (IPCC 2021). Other aspects of the climate are also changing, such as rainfall patterns, snow and ice cover, and sea level. If greenhouse gas concentrations continue to increase, climate models predict that the average temperature at the Earth's surface is likely to increase between 0.27 to 9.99 degrees Fahrenheit (0.15 to 5.55 degrees Celsius) relative to 1995 to 2014 levels by the end of this century, depending on the emissions scenario and the responsiveness of the climate system (IPCC 2021).

For further information on greenhouse gases, radiative forcing, and implications for climate change, see the recent scientific assessment reports from the IPCC,³⁵ the U.S. Global Change Research Program (USGCRP),³⁶ and the National Academies of Sciences, Engineering, and Medicine (NAS).³⁷

Greenhouse Gases

Although the Earth's atmosphere consists mainly of oxygen and nitrogen, neither plays a significant role in enhancing the greenhouse effect because both are essentially transparent to terrestrial radiation. The greenhouse effect is primarily a function of the concentration of water vapor, carbon dioxide (CO₂), methane (CH₄), nitrous oxide (N₂O), and other trace gases in the atmosphere that absorb the terrestrial radiation leaving the surface of the Earth (IPCC 2021).

Naturally occurring greenhouse gases include water vapor, CO₂, CH₄, N₂O, and ozone (O₃). Several classes of halogenated substances that contain fluorine, chlorine, or bromine are also greenhouse gases, but they are, for the most part, solely a product of industrial activities. Chlorofluorocarbons (CFCs) and hydrochlorofluorocarbons (HCFCs) are halocarbons that contain chlorine, while halocarbons that contain bromine are referred to as bromofluorocarbons (i.e., halons). As stratospheric ozone depleting substances, CFCs, HCFCs, and halons are covered under the Montreal Protocol on Substances that Deplete the Ozone Layer. The UNFCCC defers to this earlier international treaty. Consequently, Parties to the UNFCCC are not required to include these gases in

³² On October 30, 2009, EPA promulgated a rule requiring annual reporting of greenhouse gas data from large greenhouse gas emissions sources in the United States. Implementation of the rule, codified at 40 CFR Part 98, is referred to as EPA's Greenhouse Gas Reporting Program (GHGRP).

³³ See <http://www.epa.gov/ghgreporting> and <http://ghgdata.epa.gov/ghgp/main.do>.

³⁴ See <https://www.ccdsupport.com/confluence/pages/viewpage.action?pageId=322699300>

³⁵ See <https://www.ipcc.ch/report/ar6/wg1/>.

³⁶ See <https://nca2018.globalchange.gov/>.

³⁷ See <https://www.nationalacademies.org/topics/climate>.

national greenhouse gas inventories.³⁸ Some other fluorine-containing halogenated substances—hydrofluorocarbons (HFCs), perfluorocarbons (PFCs), sulfur hexafluoride (SF₆), and nitrogen trifluoride (NF₃)—do not deplete stratospheric ozone but are potent greenhouse gases. These latter substances are addressed by the UNFCCC and accounted for in national greenhouse gas inventories.

There are also several other substances that influence the global radiation budget but are short-lived and therefore not well-mixed, leading to spatially variable radiative forcing effects. These substances include carbon monoxide (CO), nitrogen dioxide (NO₂), sulfur dioxide (SO₂), and tropospheric (ground level) ozone (O₃). Tropospheric ozone is formed from chemical reactions in the atmosphere of precursor pollutants, which include volatile organic compounds (VOCs), CH₄, and nitrogen oxides (NO_x), in the presence of ultraviolet light (sunlight).

Aerosols are extremely small particles or liquid droplets suspended in the Earth’s atmosphere that are often composed of sulfur compounds, carbonaceous combustion products (e.g., black carbon), crustal materials (e.g., dust) and other human-induced pollutants. They can affect the absorptive characteristics of the atmosphere (e.g., scattering incoming sunlight away from the Earth’s surface, or, in the case of black carbon, absorbing sunlight) and can play a role in affecting cloud formation and lifetime, as well as the radiative forcing of clouds and precipitation patterns.

CO₂, CH₄, and N₂O are continuously emitted to and removed from the atmosphere by natural processes on Earth. Anthropogenic activities (such as fossil fuel combustion, cement production, land-use, land-use change, and forestry, agriculture, or waste management), however, can cause additional quantities of these and other greenhouse gases to be emitted or sequestered, thereby changing their global average atmospheric concentrations. Natural activities such as respiration by plants or animals and seasonal cycles of plant growth and decay are examples of processes that only cycle carbon or nitrogen between the atmosphere and organic biomass. Such processes, except when directly or indirectly perturbed out of equilibrium by anthropogenic activities, generally do not alter average atmospheric greenhouse gas concentrations over decadal timeframes. Climatic changes resulting from anthropogenic activities, however, could have positive or negative feedback effects on these natural systems. Atmospheric concentrations of these gases, along with their rates of growth and atmospheric lifetimes, are presented in Table 1-1.

Table 1-1: Global Atmospheric Concentration, Rate of Concentration Change, and Atmospheric Lifetime of Selected Greenhouse Gases

Atmospheric Variable	CO ₂	CH ₄	N ₂ O	SF ₆	CF ₄
Pre-industrial atmospheric concentration	280 ppm	0.730 ppm	0.270 ppm	0.01 ppt	34.1 ppt
Atmospheric concentration	419 ppm ^a	1.912 ppm ^b	0.336 ppm ^c	11.02 ppt ^d	85.5 ppt ^e
Rate of concentration change	2.28 ppm/yr ^f	8.83 ppb/yr ^{f,g}	1.01 ppb/yr ^f	0.32 ppt/yr ^f	0.81 ppt/yr ^f
Atmospheric lifetime (years)	See footnote ^h	11.8	109 ⁱ	About 1,000 ^j	50,000

^a The atmospheric CO₂ concentration is the 2022 annual average at the Mauna Loa, HI station (NOAA/ESRL 2024a). The global atmospheric CO₂ concentration, computed using an average of sampling sites across the world, was 417 ppm in 2022.

^b The values presented are global 2022 annual average mole fractions (NOAA/ESRL 2024b).

^c The values presented are global 2022 annual average mole fractions (NOAA/ESRL 2024c).

^d The values presented are global 2022 annual average mole fractions (NOAA/ESRL 2024d).

^e The 2019 CF₄ global mean atmospheric concentration is from the Advanced Global Atmospheric Gases Experiment (IPCC 2021).

^f The rate of concentration change for CO₂ is an average of the rates from 2007 through 2022 and has fluctuated between 1.5 to 3.0 ppm per year over this period (NOAA/ESRL 2024a). The rate of concentration change for CH₄, N₂O, and SF₆, is the average rate of change between 2007 and 2022 (NOAA/ESRL 2024b; NOAA/ESRL 2024c; NOAA/ESRL 2024d). The rate of concentration change for CF₄ is the average rate of change between 2011 and 2019 (IPCC 2021).

^g The growth rate for atmospheric CH₄ decreased from over 10 ppb/year in the 1980s to nearly zero in the early 2000s; recently, the growth rate has been about 13.22 ppb/year (NOAA/ESRL 2024b).

³⁸ Emissions estimates of CFCs, HCFCs, halons and other ozone-depleting substances are included in this document for informational purposes.

^h For a given amount of CO₂ emitted, some fraction of the atmospheric increase in concentration is quickly absorbed by the oceans and terrestrial vegetation, some fraction of the atmospheric increase will only slowly decrease over a number of years, and a small portion of the increase will remain for many centuries or more.

ⁱ This table reports the “perturbation lifetime” for both CH₄ and N₂O, which takes into account the interactions between emissions of the gas and its own atmospheric residence time.

^j The lifetime for SF₆ was revised from 3,200 years to about 1,000 years based on recent studies (IPCC 2021).

Source: Pre-industrial atmospheric concentrations and atmospheric lifetimes for CH₄ and N₂O, are from IPCC (2021), pre-industrial atmospheric concentration for SF₆, is from Rigby (2010), and pre-industrial atmospheric concentration for CF₄ is from Meinhausen (2017).

A brief description of each greenhouse gas, its sources, and its role in the atmosphere is given below. The following section then explains the concept of global warming potentials (GWPs), which are assigned to individual gases as a measure of their relative average global radiative forcing effect.

Water Vapor (H₂O). Water vapor is the largest contributor to the natural greenhouse effect. Water vapor is fundamentally different from other greenhouse gases in that it can condense and rain out when it reaches high concentrations, and the total amount of water vapor in the atmosphere is in part a function of the Earth’s temperature. While some human activities such as evaporation from irrigated crops or power plant cooling release water vapor into the air, these activities have been determined to have a negligible effect on global climate (IPCC 2021). The lifetime of water vapor in the troposphere is on the order of 10 days. Water vapor can also contribute to cloud formation, and clouds can have both warming and cooling effects by either trapping or reflecting heat. Because of the relationship between water vapor levels and temperature, water vapor and clouds serve as a feedback to climate change, such that for any given change in other greenhouse gas concentrations, the total temperature change is greater than would happen in the absence of water vapor. Aircraft emissions can create contrails, which may also develop into contrail-induced cirrus clouds, with complex regional and temporal net radiative forcing effects that currently have a low level of scientific certainty (IPCC 2021).

Carbon Dioxide (CO₂). In nature, carbon is cycled between various atmospheric, oceanic, land biotic, marine biotic, and mineral reservoirs. The largest fluxes occur between the atmosphere and terrestrial biota, and between the atmosphere and surface water of the oceans. In the atmosphere, carbon predominantly exists in its oxidized form as CO₂. Atmospheric CO₂ is part of this global carbon cycle, and therefore its fate is a complex function of geochemical and biological processes. Carbon dioxide concentrations in the atmosphere increased from approximately 280 parts per million by volume (ppmv) in pre-industrial times to 419 ppmv in 2022, a 50 percent increase (IPCC 2021; NOAA/ESRL 2024a).^{39, 40} The IPCC states that “Observed increases in well-mixed greenhouse gas (GHG) concentrations since around 1750 are unequivocally caused by human activities” (IPCC 2021). The predominant source of anthropogenic CO₂ emissions is the combustion of fossil fuels. Forest clearing, other biomass burning, and some non-energy production processes (e.g., cement production) also emit notable quantities of CO₂. In its *Sixth Assessment Report*, the IPCC determined that of the 2.0 degrees of observed warming, the best estimate is that 1.9 degrees of that are due to human influence, with elevated CO₂ concentrations being the most important contributor to that warming (IPCC 2021).

Methane (CH₄). Methane is primarily produced through anaerobic decomposition of organic matter in biological systems. Agricultural processes such as wetland rice cultivation, enteric fermentation in animals, and the decomposition of animal wastes emit CH₄, as does the decomposition of municipal solid wastes and treatment of wastewater. Methane is also emitted during the production and distribution of natural gas and petroleum, and is released as a byproduct of coal mining and incomplete fossil fuel combustion. Atmospheric concentrations of CH₄ have increased by about 162 percent since 1750, from a pre-industrial value of about 730 ppb to 1,912 ppb in 2022⁴¹ although the rate of increase decreased to near zero in the early 2000s, and has recently increased again to

³⁹ The pre-industrial period is considered as the time preceding the year 1750 (IPCC 2021).

⁴⁰ Carbon dioxide concentrations during the last 1,000 years of the pre-industrial period (i.e., 750 to 1750), a time of relative climate stability, fluctuated by about ±10 ppmv around 280 ppmv (IPCC 2021).

⁴¹ This value is the global 2022 annual average mole fraction (NOAA/ESRL 2024b).

about 8.83 ppb/year. The IPCC has estimated that about half of the current CH₄ flux to the atmosphere (and the entirety of the increase in concentration) is anthropogenic, from human activities such as agriculture, fossil fuel production and use, and waste disposal (IPCC 2021).

Methane is primarily removed from the atmosphere through a reaction with the hydroxyl radical (OH) and is ultimately converted to CO₂. Minor removal processes also include reaction with chlorine in the marine boundary layer, a soil sink, and stratospheric reactions. Increasing emissions of CH₄ reduce the concentration of OH, creating a feedback that increases the atmospheric lifetime of CH₄ (IPCC 2021). Methane's reactions in the atmosphere also lead to production of tropospheric ozone and stratospheric water vapor, both of which also contribute to climate change. Tropospheric ozone also has negative effects on human health and plant productivity.

Nitrous Oxide (N₂O). Anthropogenic sources of N₂O emissions include agricultural soils, especially production of nitrogen-fixing crops and forages, the use of synthetic and manure fertilizers, and manure deposition by livestock; fossil fuel combustion, especially from mobile combustion; adipic (nylon) and nitric acid production; wastewater treatment and waste incineration; and biomass burning. The atmospheric concentration of N₂O has increased by 24 percent since 1750, from a pre-industrial value of about 270 ppb to 336 ppb in 2022,⁴² a concentration that has not been exceeded during at least the last 800 thousand years. Nitrous oxide is primarily removed from the atmosphere by the photolytic action of sunlight in the stratosphere (IPCC 2021).

Ozone (O₃). Ozone is present in both the upper stratosphere,⁴³ where it shields the Earth from harmful levels of ultraviolet radiation, and at lower concentrations in the troposphere,⁴⁴ where it is the main component of anthropogenic photochemical "smog." During the last two decades, emissions of anthropogenic chlorine and bromine-containing halocarbons, such as CFCs, have depleted stratospheric ozone concentrations. This loss of ozone in the stratosphere has resulted in negative radiative forcing, representing an indirect effect of anthropogenic emissions of chlorine and bromine compounds (IPCC 2021). The depletion of stratospheric ozone and its radiative forcing remained relatively unchanged since 2000 for the last two decades and is starting to decline; recovery is expected to occur shortly after the middle of the twenty-first century (WMO/UNEP 2018).

The past increase in tropospheric ozone, which is also a greenhouse gas, is estimated to provide the third largest increase in direct radiative forcing since the pre-industrial era, behind CO₂ and CH₄. Tropospheric ozone is produced from complex chemical reactions of volatile organic compounds and CH₄ mixing with NO_x in the presence of sunlight. The tropospheric concentrations of ozone and these other pollutants are short-lived and, therefore, spatially variable (IPCC 2021).

Halocarbons, Sulfur Hexafluoride (SF₆), and Nitrogen Trifluoride (NF₃). Halocarbons are, for the most part, man-made chemicals that have direct radiative forcing effects and could also have an indirect effect. Halocarbons that contain chlorine (CFCs, HCFCs, methyl chloroform, and carbon tetrachloride) and bromine (halons, methyl bromide, and hydrobromofluorocarbons) result in stratospheric ozone depletion and are therefore controlled under the Montreal Protocol on Substances that Deplete the Ozone Layer. Although most CFCs and HCFCs are potent global warming gases, their net radiative forcing effect on the atmosphere is reduced because they cause stratospheric ozone depletion, which itself is a greenhouse gas but which also shields the Earth from harmful levels of ultraviolet radiation. Under the Montreal Protocol, the United States phased out the production and importation of halons by 1994 and of CFCs by 1996. Under the Copenhagen Amendments to the Protocol, a cap

⁴² This value is the global 2022 annual average (NOAA/ESRL 2024c).

⁴³ The stratosphere is the layer from the troposphere up to roughly 50 kilometers. In the lower regions the temperature is nearly constant but in the upper layer the temperature increases rapidly because of sunlight absorption by the ozone layer. The ozone-layer is the part of the stratosphere from 19 kilometers up to 48 kilometers where the concentration of ozone reaches up to 10 parts per million.

⁴⁴ The troposphere is the layer from the ground up to 11 kilometers near the poles and up to 16 kilometers in equatorial regions (i.e., the lowest layer of the atmosphere where people live). It contains roughly 80 percent of the mass of all gases in the atmosphere and is the site for most weather processes, including most of the water vapor and clouds.

was placed on the production and importation of HCFCs by non-Article 5 countries, including the United States,⁴⁵ beginning in 1996, and then followed by intermediate requirements and a complete phase-out by the year 2030. While ozone depleting gases covered under the Montreal Protocol and its Amendments are not covered by the UNFCCC, they are reported in this Inventory under Annex 6.2 for informational purposes.

HFCs, PFCs, SF₆, and NF₃ are not ozone depleting substances. The most common HFCs are, however, powerful greenhouse gases. Hydrofluorocarbons are primarily used as replacements for ozone depleting substances but are also emitted as a byproduct of the HCFC-22 (chlorodifluoromethane) manufacturing process. Other contributing sources to HFC emissions include the electronics industry and magnesium production and processing. Currently, these emissions have a small aggregate radiative forcing impact, but it was anticipated that without further controls their contribution to overall radiative forcing would increase, the ERF (effective radiative forcing) of halogenated gases increased by 3.5 percent between 2011 and 2019 primarily due to a decrease in atmospheric mixing-ratios of CFCs and an increase in their replacements (IPCC 2021). On December 27, 2020, the American Innovation and Manufacturing (AIM) Act was enacted by Congress and which gives EPA authority to phase down HFC production and consumption (i.e., production plus import, minus export), through an allowance allocation program, promulgate certain regulations for purposes of maximizing reclamation and minimizing releases of HFCs and their substitutes from equipment, and facilitate the transition to next-generation technologies through sector-based restrictions, which will lead to lower HFC emissions over time. On October 31, 2022, the United States also ratified the Kigali Amendment to the Montreal Protocol, committing to continued phase down of HFCs. Perfluorocarbons, SF₆, and NF₃ are predominantly emitted from various industrial processes including aluminum smelting, semiconductor manufacturing, electric power transmission and distribution, and magnesium casting. Currently, the radiative forcing impact of PFCs, SF₆, and NF₃ is also small, but they have a significant growth rate, extremely long atmospheric lifetimes, and are strong absorbers of infrared radiation, and therefore have the potential to influence climate far into the future (IPCC 2021).

Carbon Monoxide (CO). Carbon monoxide has an indirect radiative forcing effect by elevating concentrations of CH₄ and tropospheric ozone through chemical reactions with other atmospheric constituents (e.g., the hydroxyl radical, OH) that would otherwise assist in destroying CH₄ and tropospheric ozone. Carbon monoxide is created when carbon-containing fuels are burned incompletely. Through natural processes in the atmosphere, it is eventually oxidized to CO₂. Carbon monoxide concentrations are both short-lived in the atmosphere and spatially variable.

Nitrogen Oxides (NO_x). The primary climate change effects of nitrogen oxides (i.e., NO and NO₂) are indirect. Warming effects can occur due to reactions leading to the formation of ozone in the troposphere, but cooling effects can occur due to the role of NO_x as a precursor to nitrate particles (i.e., aerosols) and due to destruction of stratospheric ozone when emitted from very high-altitude aircraft.⁴⁶ Additionally, NO_x emissions are also likely to decrease CH₄ concentrations, thus having a negative radiative forcing effect (IPCC 2021). Nitrogen oxides are created from lightning, soil microbial activity, biomass burning (both natural and anthropogenic fires) fuel combustion, and, in the stratosphere, from the photo-degradation of N₂O. Concentrations of NO_x are both relatively short-lived in the atmosphere and spatially variable.

Non-methane Volatile Organic Compounds (NMVOCs). Non-methane volatile organic compounds include substances such as propane, butane, and ethane. These compounds participate, along with NO_x, in the formation of tropospheric ozone and other photochemical oxidants. NMVOCs are emitted primarily from transportation and industrial processes, as well as biomass burning and non-industrial consumption of organic solvents. Concentrations of NMVOCs tend to be both short-lived in the atmosphere and spatially variable.

⁴⁵ Article 5 of the Montreal Protocol covers several groups of countries, especially developing countries, with low consumption rates of ozone depleting substances. Developing countries with per capita consumption of less than 0.3 kg of certain ozone depleting substances (weighted by their ozone depleting potential) receive financial assistance and a grace period of ten additional years in the phase-out of ozone depleting substances.

⁴⁶ NO_x emissions injected higher in the stratosphere, primarily from fuel combustion emissions from high altitude supersonic aircraft, can lead to stratospheric ozone depletion.

Aerosols. Aerosols are extremely small particles or liquid droplets found in the atmosphere that are either directly emitted into or are created through chemical reactions in the Earth’s atmosphere. Aerosols or their chemical precursors can be emitted by natural events such as dust storms, biogenic or volcanic activity, or by anthropogenic processes such as transportation, coal combustion, cement manufacturing, waste incineration, or biomass burning. Various categories of aerosols exist from both natural and anthropogenic sources, such as soil dust, sea salt, biogenic aerosols, sulfates, nitrates, volcanic aerosols, industrial dust, and carbonaceous⁴⁷ aerosols (e.g., black carbon, organic carbon). Aerosols can be removed from the atmosphere relatively rapidly by precipitation or through more complex processes under dry conditions.

Aerosols affect radiative forcing differently than greenhouse gases. Their radiative effects occur through direct and indirect mechanisms: directly by scattering and absorbing solar radiation (and to a lesser extent scattering, absorption, and emission of terrestrial radiation); and indirectly by increasing cloud droplets and ice crystals that modify the formation, precipitation efficiency, and radiative properties of clouds (IPCC 2021). Despite advances in understanding of cloud-aerosol interactions, the contribution of aerosols to radiative forcing are difficult to quantify because aerosols generally have short atmospheric lifetimes, and have number concentrations, size distributions, and compositions that vary regionally, spatially, and temporally (IPCC 2021).

The net effect of aerosols on the Earth’s radiative forcing is believed to be negative (i.e., net cooling effect on the climate). In fact, aerosols contributed a cooling influence of up to 1.4 degrees, offsetting a substantial portion of greenhouse gas warming (IPCC 2021). Because aerosols remain in the atmosphere for only days to weeks, their concentrations respond rapidly to changes in emissions.⁴⁸ Not all aerosols have a cooling effect. Current research suggests that another constituent of aerosols, black carbon, has a positive radiative forcing by heating the Earth’s atmosphere and causing surface warming when deposited on ice and snow (IPCC 2021). Black carbon also influences cloud development, but the direction and magnitude of this forcing is an area of active research.

Global Warming Potentials

A GWP is a quantified measure of the relative globally averaged radiative forcing impacts of emissions of a particular greenhouse gas over time (see Table 1-2). It is defined as the accumulated radiative forcing within a specific time horizon caused by emitting 1 kilogram (kg) of the gas, relative to that of the reference gas CO₂ (IPCC 2021). Direct radiative effects occur when the gas itself absorbs radiation. Indirect radiative forcing occurs when chemical transformations involving the original gas produce a gas or gases that are greenhouse gases, or when a gas influences other radiatively important processes such as the atmospheric lifetimes of other gases. The reference gas used is CO₂, and therefore GWP-weighted emissions are measured in CO₂ equivalent (CO₂ Eq.).⁴⁹ For example, the relationship between a kg of emissions of a gas and a kg of CO₂ Eq. emissions can be expressed as follows and also adapted to other units (e.g. metric tons, etc.):

⁴⁷ Carbonaceous aerosols are aerosols that are comprised mainly of carbon and hydrogen. Those carbonaceous aerosols with more hydrogen are classified as “organic carbon”, and are generally reflective, while the aerosols that are nearly pure carbon are classified as “black carbon” (also referred to as “soot”) and can absorb light (IPCC 2021).

⁴⁸ Volcanic activity can inject significant quantities of aerosol producing sulfur dioxide and other sulfur compounds into the stratosphere, which can result in a longer lasting negative forcing effect (i.e., a few years) (IPCC 2021).

⁴⁹ Carbon comprises 12/44^{ths} of carbon dioxide by weight.

Equation 1-1: Calculating CO₂ Equivalent Emissions

$$\text{kg CO}_2 \text{ Eq.} = (\text{kg emission of gas}) \times (\text{GWP})$$

where,

kg CO ₂ Eq.	= kilograms of CO ₂ equivalent
kt	= kilograms (equivalent to a thousand metric grams)
GWP	= Global warming potential

GWP values allow for a comparison of the impacts of emissions and reductions of different gases. According to the IPCC, GWPs typically have an uncertainty of ±40 percent.

All estimates are provided throughout the report in both MMT CO₂ equivalents and unweighted units. Recent decisions under the UNFCCC require Parties to use 100-year GWP values from the IPCC *Fifth Assessment Report* (AR5) for calculating CO₂-equivalent emissions in their national reporting by the end of 2024.

...Decides that, until it adopts a further decision on the matter, the global warming potential values used by Parties in their reporting under the Convention to calculate the carbon dioxide equivalence of anthropogenic greenhouse gas emissions by sources and removals by sinks shall be based on the effects of greenhouse gases over a 100-year time horizon as listed in table 8.A.1 in appendix 8.A to the contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change,⁵⁰ excluding the value for fossil methane.⁵¹

This reflects updated science and ensures that national greenhouse gas inventories reported by all nations are comparable. In preparation for upcoming UNFCCC requirement,⁵² this report reflects CO₂-equivalent greenhouse gas totals using 100-year AR5 GWP values. A comparison of emission values with the previously used 100-year GWP values from IPCC *Fourth Assessment Report* (AR4) (IPCC 2007), and the IPCC *Sixth Assessment Report* (AR6) (IPCC 2021) values can be found in Annex 6.1 of this report. The 100-year GWP values used in this report are listed below in Table 1-2.

Greenhouse gases with relatively long atmospheric lifetimes (e.g., CO₂, CH₄, N₂O, HFCs, PFCs, SF₆, NF₃) tend to be evenly distributed throughout the atmosphere, and consequently global average concentrations can be determined. The short-lived gases such as water vapor, carbon monoxide, tropospheric ozone, ozone precursors (e.g., NO_x, and NMVOCs), and tropospheric aerosols (e.g., SO₂ products and carbonaceous particles), however, vary regionally, and consequently it is difficult to quantify their global radiative forcing impacts. Parties to the UNFCCC have not agreed upon GWP values for these gases that are short-lived and spatially inhomogeneous in the atmosphere.

Table 1-2: Global Warming Potentials and Atmospheric Lifetimes (Years) Used in this Report

Gas	Atmospheric Lifetime	GWP ^a
CO ₂	See footnote ^b	1
CH ₄ ^c	12.4	28
N ₂ O	121	265

⁵⁰ Intergovernmental Panel on Climate Change. 2013. *Climate Change 2013: The Physical Science Basis*. Contribution of Working Group I to the *Fifth Assessment Report* of the Intergovernmental Panel on Climate Change. TF Stocker, D Qin, G-K Plattner, et al. (eds.). Cambridge and New York: Cambridge University Press. Available at <http://www.ipcc.ch/report/ar5/wg1>.

⁵¹ See paragraphs 1 and 2 of the decision on common metrics adopted at the 27th UNFCCC Conference of Parties (COP27), available online at https://unfccc.int/sites/default/files/resource/cp2022_10a01_E.pdf.

⁵² See Annex to decision 18/CMA.1, available online at https://unfccc.int/sites/default/files/resource/CMA2018_03a02E.pdf. The Paris Agreement reporting guidelines also clarified use of the 100-year GWPs listed in table 8.A.1 in Annex 8.A of Chapter 8 of the *Fifth Assessment Report* (AR5) of the Intergovernmental Panel on Climate Change, excluding the value for fossil methane. United Nations Framework Convention on Climate Change, see paragraph 25 of Decision 5/CMA.3 available online at https://unfccc.int/sites/default/files/resource/CMA2021_L10a2E.pdf.

HFC-23	222	12,400
HFC-32	5.2	677
HFC-41 ^d	2.8	116
HFC-125	28.2	3,170
HFC-134a	13.4	1,300
HFC-143a	47.1	4,800
HFC-152a	1.5	138
HFC-227ea	38.9	3,350
HFC-236fa	242	8,060
CF ₄	50,000	6,630
C ₂ F ₆	10,000	11,100
C ₃ F ₈	2,600	8,900
c-C ₄ F ₈	3,200	9,540
SF ₆	3,200	23,500
NF ₃	500	16,100
Other Fluorinated Gases		See Annex 6

^a 100-year time horizon.

^b For a given amount of CO₂ emitted, some fraction of the atmospheric increase in concentration is quickly absorbed by the oceans and terrestrial vegetation, some fraction of the atmospheric increase will only slowly decrease over a number of years, and a small portion of the increase will remain for many centuries or more.

^c The GWP of CH₄ includes the direct effects and those indirect effects due to the production of tropospheric ozone and stratospheric water vapor. The indirect effect due to the production of CO₂ is not included.

^d See Table A-1 of 40 CFR Part 98

Source: IPCC (2013).

Box 1-2: The IPCC Sixth Assessment Report and Global Warming Potentials

In 2021, the IPCC published its *Sixth Assessment Report* (AR6), which updated its comprehensive scientific assessment of climate change. Within the AR6 report, the GWP values of gases were revised relative to previous IPCC assessment reports, e.g., the IPCC *Fifth Assessment Report* (AR5) (IPCC 2014). Although the AR5 GWP values are used throughout this report, consistent with UNFCCC reporting requirements, it is straight-forward to review the changes to the GWP values and their impact on estimates of the total GWP-weighted emissions of the United States. In the AR6, the IPCC used more recent estimates of the atmospheric lifetimes and radiative efficiencies of some gases and updated background concentrations. The AR6 now includes climate-carbon feedback effects for non-CO₂ gases, improving the consistency between treatment of CO₂ and non-CO₂ gases. Indirect effects of gases on other atmospheric constituents (such as the effect of methane on ozone) have also been updated to match more recent science.

Table 1-3 presents the new GWP values, relative to those presented in the AR5, using the 100-year time horizon common to Paris Agreement and UNFCCC reporting.⁵³ Updated reporting guidelines under the Paris Agreement require the United States and other countries to shift to use of the IPCC *Fifth Assessment Report* (AR5) (IPCC 2013) 100-year GWP values (without feedbacks) for national inventory reporting.⁵⁴ All estimates provided throughout this report are also presented in unweighted units. For informational purposes, emission estimates that use 100-year GWPs from other recent IPCC Assessment Reports are presented in detail in Annex 6.1 of this report.

⁵³ See [Decision 7/CP.27 included in https://unfccc.int/sites/default/files/resource/cp2022_10a01_E.pdf](https://unfccc.int/sites/default/files/resource/cp2022_10a01_E.pdf).

⁵⁴ See <https://unfccc.int/process-and-meetings/transparency-and-reporting/reporting-and-review-under-the-paris-agreement>.

Table 1-3: Comparison of 100-Year GWP values

100-Year GWP Values				Comparisons to AR5	
Gas	AR5 ^a	AR5 with feedbacks ^b	AR6 ^c	AR5 with feedbacks ^b	AR6 ^c
CO ₂	1	1	1	NC	NC
CH ₄ ^d	28	34	27	6	(1)
N ₂ O	265	298	273	33	8
HFC-23	12,400	13,856	14,600	1,456	2,200
HFC-32	677	817	771	140	94
HFC-41	116	141	135	25	19
HFC-125	3,170	3,691	3,740	521	570
HFC-134a	1,300	1,549	1,530	249	230
HFC-143a	4,800	5,508	5,810	708	1,010
HFC-152a	138	167	164	29	26
HFC-227ea	3,350	3,860	3,600	510	250
HFC-236fa	8,060	8,998	8,690	938	630
CF ₄	6,630	7,349	7,380	719	750
C ₂ F ₆	11,100	12,340	12,400	1,240	1,300
C ₃ F ₈	8,900	9,878	9,290	978	390
c-C ₄ F ₈	9,540	10,592	10,200	1,052	660
SF ₆	23,500	26,087	24,300	2,587	800
NF ₃	16,100	17,885	17,400	1,785	1,300

NC (No Change)

^a The GWP values in this column reflect values used in this report from AR5 excluding climate-carbon feedbacks and the value for fossil methane.

^b The GWP values in this column are from the AR5 report but include climate-carbon feedbacks for the non-CO₂ gases in order to be consistent with the approach used in calculating the CO₂ lifetime.

^c The GWP values in this column are from the AR6 report.

^d The GWP of CH₄ includes the direct effects and those indirect effects due to the production of tropospheric ozone and stratospheric water vapor. Including the indirect effect due to the production of CO₂ resulting from methane oxidation would lead to an increase in AR5 methane GWP values by 2 for fossil methane and is not shown in this table.

Note: Parentheses indicate negative values.

Sources: IPCC (2021), IPCC (2013), IPCC (2007), IPCC (2001), IPCC (1996).

1.2 National Inventory Arrangements

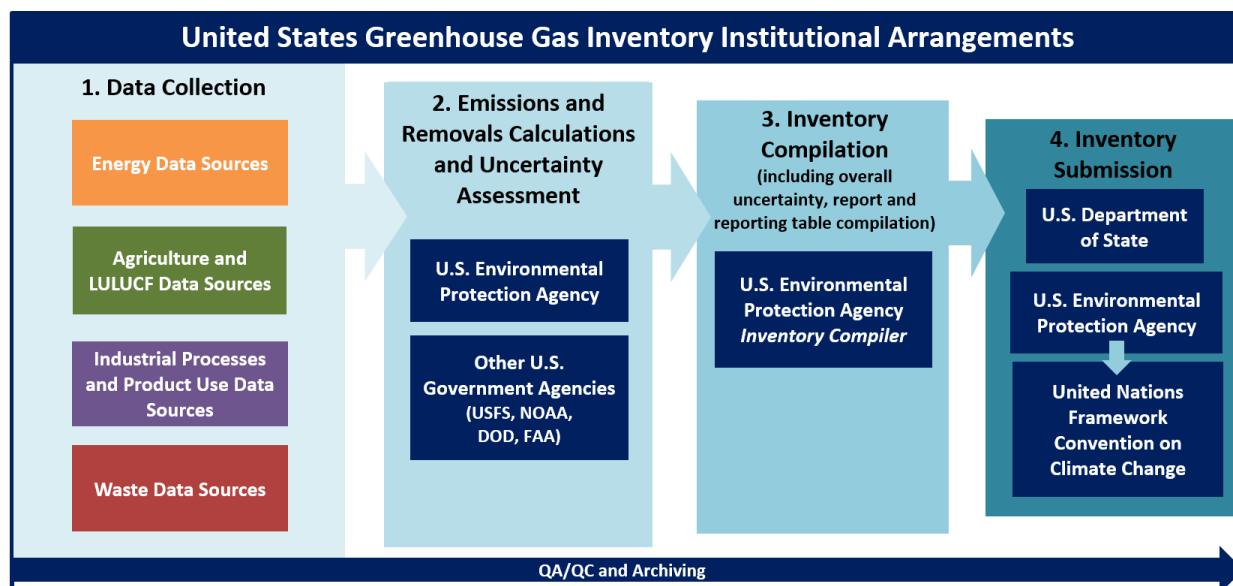
The U.S. Environmental Protection Agency (EPA), in cooperation with other U.S. government agencies, prepares the *Inventory of U.S. Greenhouse Gas Emissions and Sinks*. A wide range of agencies and individuals are involved in supplying data to, planning methodological approaches and improvements, reviewing, or preparing portions of the Inventory—including federal and state government authorities, research and academic institutions, industry associations, and private consultants.

Within EPA, the Office of Atmospheric Protection (OAP) is the lead office responsible for the emission and removal calculations provided in the Inventory, as well as the completion of the National Inventory Report including the Common Reporting Tables (CRTs). EPA's Office of Transportation and Air Quality (OTAQ) and Office of Research and Development (ORD) are also involved in calculating emissions and removals for the Inventory. The U.S. Department of State (DOS) serves as the overall national focal point to the Paris Agreement and the UNFCCC, and EPA's OAP serves as the National Inventory Focal Point for this report, including responding to technical questions and comments on the U.S. Inventory. EPA staff coordinate the annual methodological choice, activity data collection, emission and removal calculations, uncertainty assessment, QA/QC processes, and improvement

planning at the individual source and sink category level. EPA’s inventory coordinator manages overall compilation of the entire *Inventory* into the proper reporting format for submission under the Paris Agreement and the UNFCCC and is responsible for the synthesis of information along with the consistent application of cross-cutting IPCC good practice across the *Inventory*.

Several other government agencies contribute to the collection and analysis of the necessary underlying activity data used in the *Inventory* calculations via formal (e.g., interagency agreements) and informal relationships, in addition to the calculation of estimates integrated in the report (e.g., U.S. Department of Agriculture’s U.S. Forest Service and Agricultural Service, National Oceanic and Atmospheric Administration, Federal Aviation Administration, and Department of Defense). Other U.S. agencies provide official data for use in the *Inventory*. The U.S. Department of Energy’s Energy Information Administration provides national fuel consumption data and the U.S. Department of Defense provides data on military fuel consumption and use of bunker fuels. Other U.S. agencies providing activity data for use in EPA’s emission calculations include: the U.S. Department of Agriculture, National Oceanic and Atmospheric Administration, the U.S. Geological Survey, the Federal Highway Administration, the Department of Transportation, the Bureau of Transportation Statistics, the Department of Commerce, and the Federal Aviation Administration. Academic and research centers also provide activity data and calculations to EPA, as well as individual companies participating in voluntary outreach efforts with EPA. EPA engages with agencies regularly on data needs and improvements to ensure sufficient activity collection for annual compilation of estimates. Finally, EPA as the National Inventory Focal Point, in coordination with the U.S. Department of State, officially submits the *Inventory* under the Paris Agreement and the UNFCCC each April by the reporting deadline.

Figure 1-1: National Inventory Arrangements and Process Diagram



Overview of Inventory Data Sources by Source and Sink Category

Energy	Agriculture and LULUCF	IPPU	Waste
U.S. Energy Information Administration	USDA U.S. Forest Service Forest Inventory and Analysis Program (FIA)	EPA Greenhouse Gas Reporting Program (GHGRP)	EPA Greenhouse Gas Reporting Program (GHGRP)
U.S. Department of Commerce – Bureau of the Census	USDA Natural Resource Conservation Service (NRCS)	U.S. Geological Survey (USGS) National Minerals Information Center	EPA Office of Land and Emergency Management (OLEM)
U.S. Department of Defense – Defense Logistics Agency	USDA National Agricultural Statistics Service (NASS) and Agricultural Research Service (ARS)	American Chemistry Council (ACC)	EPA Clean Watershed Needs Survey (CWNS)
U.S. Department of Homeland Security	EPA Office of Research and Development (ORD)	American Iron and Steel Institute (AISI)	American Housing Survey
U.S. Department of Transportation - Federal Highway Administration	U.S. Fish and Wildlife Service	U.S. International Trade Commission (USITC)	Data from research studies, trade publications, and industry associations
U.S. Department of Transportation - Federal Aviation Administration	U.S. Department of Agriculture (USDA) Animal and Plant Health Inspection Service (APHIS)	Air-Conditioning, Heating, and Refrigeration Institute	
U.S. Department of Transportation & Bureau of Transportation Statistics	Association of American Plant Food Control Officials (AAPFCO)	Data from other U.S. government agencies, research studies, trade publications, and industry association	
U.S. Department of Labor – Mine Safety and Health Administration	National Oceanic and Atmospheric Administration (NOAA)	UNEP Technology and Economic Assessment Panel (TEAP)	
U.S. Department of Energy and its National Laboratories	EPA Office of Land and Emergency Management (OLEM)		
EPA Acid Rain Program	USDA Farm Service Agency		
EPA MOVES Model	U.S. Geological Survey (USGS)		
EPA Greenhouse Gas Reporting Program (GHGRP)	U.S. Department of the Interior (DOI) - Bureau of Land Management (BLM)		
U.S. Department of Labor – Mine Safety and Health Administration	EPA Office of Land and Emergency Management (OLEM)		
American Association of Railroads	Alaska Department of Natural Resources		
American Public Transportation Association	U.S. Department of Commerce – Bureau of the Census		
U.S. Department of Interior - Bureau of Ocean Energy Management	Data from research studies, trade publications, and industry associations		
Federal Energy Regulatory Commission			
Data from research studies, trade publications, and industry associations			

Note: This table is not an exhaustive list of all data sources.

1.3 Inventory Preparation Process

This section describes EPA’s approach to preparing the annual U.S. *Inventory*, which includes both the National Inventory Document (NID) and Common Reporting Tables (CRTs). The inventory coordinator at EPA, with support from the cross-cutting compilation staff, is responsible for coordinating aggregation of all emission and removal estimates, conducting the overall uncertainty analysis of *Inventory* emissions and trends over time, and ensuring consistency and quality throughout the NID and CRTs. Emission and removal calculations, including associated uncertainty analysis for individual sources and/or sink categories are the responsibility of individual source and sink category leads, who are most familiar with each category, underlying data, and the unique national circumstances relevant to its emissions or removals profile. Using IPCC methodological decision trees and suggested good practice guidance, the individual leads determine the most appropriate methodology and collect the relevant activity data to use in the emission and removal calculations, based upon their expertise in the source or sink category, as well as coordinating with researchers and expert consultants familiar with the sources and sinks. Each year, the coordinator oversees a multi-stage process for collecting information from each individual source and sink category lead to compile all information and data for the *Inventory*.

Methodology Development, Data Collection, and Emissions and Sinks Estimation

Source and sink category leads at EPA coordinate the collection of input data (e.g., activity data and other information) and, as necessary, evaluate or develop the estimation methodology for the individual source and/or sink categories. Because EPA has been leading preparation of the *Inventory* for many years, for most source and sink categories, the methodology for the previous year is applied to the new “current” year of the *Inventory*, and inventory analysts collect any new data or update data that have changed from the previous year. If estimates for a new source or sink category are being developed for the first time, or if the methodology is changing for an existing category (e.g., the United States is implementing improvement efforts to apply a higher tiered approach for that category), then the source and/or sink category lead will develop and implement the new or refined methodology, gather the appropriate activity data and other information (e.g., emission factors or in some cases direct emission measurements) for the entire time series, and conduct any further category-specific review with involvement of relevant experts from industry, government, and universities (see Chapter 9 and Box ES-3 on EPA’s approach to recalculations).

Once the methodology is in place and the data are collected, the individual source and sink category leads calculate emission and removal estimates. The individual leads then update or create the relevant national inventory document text and accompanying annexes for the *Inventory*. Source and sink category leads are also responsible for completing the relevant sectoral background tables of the CRTs, conducting quality control (QC) checks, preparing relevant category materials for QA, or expert reviews, category-level uncertainty assessments, and reviewing data for publication in EPA’s GHG Data Explorer.⁵⁵

The treatment of confidential business information (CBI) in the *Inventory* is based on EPA internal guidelines, as well as regulations⁵⁶ applicable to the data used. EPA has specific procedures in place to safeguard CBI during the inventory compilation process. When information derived from CBI data is used for development of inventory calculations, EPA procedures ensure that these confidential data are sufficiently aggregated to protect confidentiality while still providing useful information for analysis. For example, within the Energy and Industrial

⁵⁵ See <https://cfpub.epa.gov/ghgdata/inventoryexplorer/>.

⁵⁶ 40 CFR Part 2, Subpart B titled “Confidentiality of Business Information” which is the regulation establishing rules governing handling of data entitled to confidentiality treatment. See <https://www.ecfr.gov/cgi-bin/text-idx?SID=a764235c9eadf9afe05fe04c07a28939&mc=true&node=sp40.1.2.b&rgn=div6>.

Processes and Product Use (IPPU) sectors, EPA has used aggregated facility-level data from the Greenhouse Gas Reporting Program (GHGRP) to develop, inform, and/or quality-assure U.S. emission estimates. In 2014, EPA's GHGRP, with industry engagement, compiled criteria that would be used for aggregating its confidential data to shield the underlying CBI from public disclosure.⁵⁷ In the *Inventory*, EPA is publishing only data values that meet the GHGRP aggregation criteria.⁵⁸ Specific uses of aggregated facility-level data are described in the respective methodological sections within those chapters. In addition, EPA uses historical data reported voluntarily to EPA via various voluntary initiatives with U.S. industry (e.g., EPA Voluntary Aluminum Industrial Partnership (VAIP)) and follows guidelines established under the voluntary programs for managing CBI.

Data Compilation and Archiving

The inventory coordinator at EPA with support from the data/document manager collects the source and sink categories' descriptive text and annexes, and also aggregates the emission and removal estimates into a summary data file that links the individual source and sink category data files together. This summary data file contains all of the essential data in one central location, in formats commonly used in the *Inventory* document. In addition to the data from each source and sink category, other national trend and related data are also gathered in the summary sheet for use in the Executive Summary, Introduction, and Trends chapters of the *Inventory* report (e.g., GDP, population, energy use). Similarly, the recalculation analysis and key category analysis are completed in a separate data file based on output from the summary data file. The uncertainty estimates for each source and sink category are also aggregated into uncertainty summary data files that are used to conduct the overall *Inventory* uncertainty analysis (see Section 1.7). A Microsoft SharePoint work site, maintained within EPA's IT infrastructure by the inventory coordinator, provides a platform for facilitating collaboration on the national inventory report preparation during each compilation phase, but also the efficient storage and archiving of electronic document and data files each annual cycle. Previous final published inventories are also maintained on a report archive page on EPA's Greenhouse Gas Emissions website.⁵⁹

National Inventory Document (NID) Preparation

The NID is compiled from the sections developed by each individual source or sink category lead. In addition, the inventory coordinator prepares a brief overview of each chapter that summarizes the emissions and removals from all sources and sinks discussed in the chapters. Also at this time, the Executive Summary, Introduction, Trends in Greenhouse Gas Emissions and Removals, and Recalculations and Improvements chapters are drafted, to reflect the trends and impact from improvements for the time series of the current *Inventory*. The analysis of trends necessitates gathering supplemental data, including annual climate, economic activity and gross domestic product, population, atmospheric conditions, and the annual use of electricity, energy, fossil and non-fossil fuels. Changes in these data are used to explain the trends observed in greenhouse gas emissions in the United States. Furthermore, specific factors that affect individual sectors are researched and discussed. Many of the factors that affect emissions are included in the *Inventory* document as separate analyses or side discussions in boxes within the text. Finally, the uncertainty analysis and key category analysis are compiled and updated in the report as part of final analysis steps. Throughout the report text boxes are also created to provide additional documentation (e.g., definitions) and/or examine the data aggregated in different ways than in the remainder of the document, such as a focus on transportation activities or emissions from electricity generation. The document is prepared to align with the Paris Agreement and UNFCCC reporting guidelines for National Inventory Reports while also reflecting national circumstances.

⁵⁷ Federal Register Notice on "Greenhouse Gas Reporting Program: Publication of Aggregated Greenhouse Gas Data." See pp. 79 and 110 of notice at <https://www.gpo.gov/fdsys/pkg/FR-2014-06-09/pdf/2014-13425.pdf>.

⁵⁸ U.S. EPA Greenhouse Gas Reporting Program. Developments on Publication of Aggregated Greenhouse Gas Data, November 25, 2014. See <http://www.epa.gov/ghgreporting/confidential-business-information-ghg-reporting>.

⁵⁹ See <https://www.epa.gov/ghgemissions/us-greenhouse-gas-inventory-report-archive>.

Common Reporting Tables (CRTs) Compilation

The CRTs are compiled from individual time series input data sheets completed by each individual source or sink category lead, which contain emissions and/or removals and activity data, estimates, methodological and completeness notations and associated explanations. The inventory coordinator and cross-cutting compilation staff import the U.S. category and subcategory background data into the UNFCCC's Enhanced Transparency Framework Reporting Tools to export CRTs, assuring consistency and completeness across all sectoral tables. The summary reports for emissions and removals, methods, and emission factors used, the summary tables indicating completeness of estimates (i.e., notation key NE/IE tables), the recalculation tables, and the emission and removal trends tables are automatically compiled by the online reporting software and reviewed by the inventory coordinator with support from the cross-cutting compilation staff. Internal automated quality checks within the software, as well as checks by the cross-cutting and category leads, are completed for the entire time series of CRTs before submission.

QA/QC and Uncertainty

Quality assurance and quality control (QA/QC) and uncertainty analyses are guided by the QA/QC and inventory coordinators, who help maintain the QA/QC plan and the overall uncertainty analysis procedures (see sections on QA/QC and Uncertainty, below) in collaboration with the broader inventory compilation team. The QA/QC coordinator works closely with the Inventory coordinator and source and sink category leads to ensure that a consistent QA/QC plan is implemented across all inventory categories. Similarly, the inventory coordinator ensures the uncertainty analysis is implemented consistently across all categories. The inventory QA/QC plan, outlined in Section 1.6 and Annex 8, is consistent with the quality assurance procedures outlined by EPA and IPCC good practices. The QA/QC and uncertainty findings also inform overall improvement planning, and specific improvements are noted in the Planned Improvements sections of respective categories. QA processes are outlined below.

Expert, Public, and UNFCCC/Paris Reviews

The compilation of the *Inventory* includes a two-stage review or QA process, in addition to international technical expert review following submission of the report under the UNFCCC and Paris Agreement. EPA publishes responses to comments received during both expert and public reviews with the publication of the final report on its website.⁶⁰ Responses to UNFCCC and Paris reviews are included in Annex 8 of this document.

During the first stage of review, i.e., the 30-day expert review period, a first draft of updated sectoral chapters are sent to technical experts who are not directly involved in preparing estimates. The purpose of the expert review is to provide an objective review of the methodological approaches and data sources used in the current *Inventory*, especially for sources and sinks which have experienced any changes since the previous *Inventory*. Expert review follows good practices from EPA's Peer Review handbook, i.e., the review is organized by sector, and reviewers are provided a guidance memo and charge questions to facilitate their review.⁶¹ Expert reviewers include other federal agency staff, researchers, industry experts, and others who have technical knowledge of the data, industry, and methods. EPA reviews and updates expert participation and outreach on an annual basis prior to each expert review cycle. Experts are identified in various ways; for example, many reach out to EPA with technical feedback and are added to the expert reviewer list. Reviewers are also identified through direct outreach by inventory staff based on expertise. Currently, EPA's expert list includes nearly 300 experts across all sectors. Once comments are received, they are reviewed by the source or sink lead and addressed in several ways. For example, comments suggesting methodological clarifications may be incorporated into methodological discussions prior to the next

⁶⁰ See <https://www.epa.gov/ghgemissions/draft-inventory-us-greenhouse-gas-emissions-and-sinks-1990-2022>.

⁶¹ See <https://www.epa.gov/osa/peer-review-handbook-4th-edition-2015>.

review phase, while comments citing new literature or data will be noted for review as part of planned improvements.

Following expert review, a second draft of the document, including cross-cutting synthesis chapters, is released for a 30-day public review through a notice in the U.S. Federal Register. The entire draft *Inventory* document is published on the EPA website. The public review period is open to the entire U.S. public. Comments are submitted and tracked using an online electronic docket that is accessible to the general public as well. Similar to expert review, some comments received may require further discussion with commenters, other experts and/or additional research. Specific *Inventory* improvements requiring further analysis as a result of comments are noted in the relevant category's Planned Improvement section.

As mentioned above, following completion and submission of the report under the UNFCCC and the Paris Agreement, the report also undergoes review by an international team of independent experts for adherence to UNFCCC/Paris reporting guidelines and consistency with IPCC methodological guidance.⁶² Feedback from all review processes that contribute to improving inventory quality over time are described within each planned improvement section and further in Annex 8. See also the Improvement Planning process discussed below.

Final Submittal and Publication under the Paris Agreement and the UNFCCC

After the final revisions to incorporate any comments from the Expert Review and Public Review periods, EPA prepares the final NIR, which includes the NID and the accompanying CRTs for electronic reporting. Prior to submission, EPA's Office of Atmospheric Protection briefs senior leadership on reporting findings and improvements since the previous report, along with an overview of feedback from the expert and public review processes.

EPA, as the National Inventory Focal Point, sends the official submission of the U.S. *Inventory* under the Paris Agreement and the UNFCCC using the UN's reporting software, coordinating with the U.S. Department of State, the overall UNFCCC focal point. Concurrently, the report is also published on EPA's website.⁶³ On EPA's website, users can also visualize and download the current time-series estimates from the GHG Inventory Data Explorer Tool,⁶⁴ and also download more detailed data presented in tables within the report and report annex in CSV format.

Improvement Planning

Each year, several emission and sink estimates in the *Inventory of U.S. Greenhouse Gas Emissions and Sinks* are recalculated and revised, through the use of better methods and/or data with the goal of improving inventory quality and reducing uncertainties, including the transparency, completeness, consistency, and overall usefulness of the report. In this effort, the United States follows the *2006 IPCC Guidelines* (IPCC 2006) and its *2019 Refinement*, which state, "Both methodological changes and refinements over time are an essential part of improving inventory quality. It is *good practice* to change or refine methods when available data have changed; the previously used method is not consistent with the IPCC guidelines for that category; a category has become key; the previously used method is insufficient to reflect mitigation activities in a transparent manner; the capacity for inventory preparation has increased; improved inventory methods become available; and/or for correction of errors." The EPA's OAP coordinates improvement planning across all sectors and also cross-cutting analyses based on annual review and input from the technical teams leading compilation of each sector's estimates, including

⁶² See <https://unfccc.int/process-and-meetings/transparency-and-reporting/reporting-and-review-under-the-convention/greenhouse-gas-inventories-annex-i-parties/review-process>.

⁶³ See <https://www.epa.gov/ghgemissions/inventory-us-greenhouse-gas-emissions-and-sinks>.

⁶⁴ See <https://cfpub.epa.gov/ghgdata/inventoryexplorer/>.

continuous improvements to the overall data and document compilation and QA/QC processes. Planned improvements are identified through QA/QC processes (including completeness checks), the key category analysis, and the uncertainty analysis. The inventory coordinator, with input from EPA source and sink category leads, maintains a log of all planned improvements, by sector and cross-cutting, tracking the category significance, specific category improvement, prioritization, anticipated time frame for implementation of each proposed improvement, and status of progress in implementing improvement. Improvements for significant or key categories are usually prioritized across all improvements unless effort would require disproportionate levels of effort and resources relative to improvements for other key categories to address.

1.4 Methodology and Data Sources

Emissions and removals of greenhouse gases from various source and sink categories have been estimated using methodologies that are consistent with the *2006 IPCC Guidelines for National Greenhouse Gas Inventories* (IPCC 2006) and its supplements and refinements. To a great extent, this report makes use of published official economic and physical statistics for activity data, emission factors and other key parameters as inputs to the methods applied. Depending on the category, activity data can include fuel consumption or deliveries, vehicle-miles traveled, raw material processed, or commodity produced, etc. Emission factors are factors that relate quantities of emissions to an activity. For more information on data sources see Section 1.2 above, Box 1-1 on use of GHGRP data, and categories' methodology sections for more information on other data sources. In addition to official statistics, the report utilizes findings from academic studies, trade association surveys and statistical reports, along with expert judgment, consistent with the *2006 IPCC Guidelines*.

The methodologies provided in the *2006 IPCC Guidelines* represent foundational methodologies for a variety of source and sink categories, and many of these methodologies continue to be improved and refined as new research and data become available. This report uses those IPCC methodologies when applicable, and supplements them with refined guidance, other available country-specific methodologies and data where possible (e.g., EPA's GHGRP). For example, as noted earlier in this chapter, this report does apply recent supplements and refinements to *2006 IPCC Guidelines* in estimating emissions and removals from coal mining, wastewater treatment and discharge, low voltage anode effects (LVAE) during aluminum production, drained organic soils, and management of wetlands, including flooded lands. Choices made regarding the methodologies and data sources used are provided in the Methodology and Time-Series Consistency discussion of each category within each sectoral chapter of the report, applying higher tiered methods when feasible, especially for key categories consistent with methodological decision trees. Where additional detail is helpful and necessary to explain methodologies and data sources used to estimate emissions, complete documentation is provided in the annexes as indicated in the methodology sections of those respective source categories (e.g., Annex 3.13 for forest land remaining forest land and land converted to forest land). Methods used for key categories (discussed below) are summarized in Annex 1.

1.5 Key Categories

The *2006 IPCC Guidelines* (IPCC 2006) and *2019 Refinement to the 2006 IPCC Guidelines* (IPCC 2019) define key categories as “inventory categories which individually, or as a group of categories (for which a common method, emission factor and activity data are applied) are prioritized within the national inventory system because their estimates have a significant influence on a country's total inventory of greenhouse gases in terms of the absolute level, the trend, or the level of uncertainty in emissions or removals. Whenever the term *key category* is used, it

includes both source and sink categories.”⁶⁵ A key category analysis identifies source or sink categories for focusing efforts to improve overall Inventory quality, including additional review when feasible.

The 2006 IPCC Guidelines (IPCC 2006) and its 2019 Refinement (2019) define several approaches, both quantitative and qualitative, to conduct a key category analysis and identify key categories both in terms of absolute level and trend, along with consideration of uncertainty. This report employs all approaches to identify key categories for the United States. The first approach, Approach 1, identifies significant or key categories without considering uncertainty in its calculations. An Approach 1 level assessment identifies all source and sink categories that cumulatively account for 95 percent of total level, i.e., total emissions (gross) in a given year when assessed in descending order of absolute magnitude. The level analysis was performed twice, including and excluding sources and sinks from the land use, land-use change, and forestry (LULUCF) sector categories. Similarly, an Approach 1 trend analysis can identify categories with trends that significantly influence overall trends by identifying all source and sink categories that cumulatively account for 95 percent of the sum all the trend assessments (e.g., percent change relative to national trend) when sorted in descending order of absolute magnitude.

The next method, Approach 2, was then implemented to identify any additional key categories not already identified from the Approach 1 level and trend assessments by considering uncertainty. The Approach 2 analysis differs from Approach 1 by incorporating each category’s uncertainty assessments in its calculations and was also performed twice, including and excluding LULUCF categories. An Approach 2 level assessment identifies all sources and sink categories that cumulatively account for 90 percent of the sum of all level assessments when sorted in descending order of magnitude. Similarly, an Approach 2 trend analysis can identify categories whose trends contribute significantly to overall trends weighing the relative trend difference with the category’s relative uncertainty assessment for 2022.

For 2022, based on the key category analysis, excluding the LULUCF sector and uncertainty, 34 categories accounted for 95 percent of emissions. Four categories account for 55 percent of emissions: CO₂ from road transport-related fuel combustion, CO₂ from coal-fired electricity generation, CO₂ from gas fired electricity generation, and CO₂ from gas-fired industrial processes. When considering uncertainties, additional categories such as emissions from substitutes for ozone depleting substances in aerosols were also identified as a key category. In the trend analysis, 32 categories were identified as key categories, and when considering uncertainties, 7 additional categories were identified as key. The trend analysis shows that CO₂ emissions from coal-fired electricity generation, in addition to CO₂ from gas fired electricity generation, CO₂ from road transport related combustion, and HFC and PFC emissions from substitutes for ozone depleting substances in the refrigeration and air conditioning sector are also significant with respect to trends over the time series.

When considering the contribution of the LULUCF sector to 2022 emissions and removals, 42 categories accounted for 95 percent of emissions and sinks, with the most significant category from LULUCF being net CO₂ emission from forest land remaining forest land. When considering uncertainties and the contribution of the LULUCF sector, additional categories such as net CO₂ emissions from grassland remaining grassland were also identified as a key category. In the trend analysis, 40 categories were identified as key, and when considering uncertainties, 2 additional categories were identified as key.

Finally, in addition to conducting Approach 1 and 2 level and trend assessments as described above, a qualitative assessment of the source and sinks categories was conducted to capture any additional key categories that were not identified using the previously described quantitative approaches. For this *Inventory*, no additional categories were identified using qualitative criteria recommend by IPCC, but EPA continues to review its qualitative assessment on an annual basis. Find more information on the key category analysis, including the approach to disaggregation of inventory estimates, see Annex 1 to this report.

⁶⁵ See Chapter 4 Volume 1, “Methodological Choice and Identification of Key Categories” in IPCC (2006) and IPCC (2019). See <http://www.ipcc-nggip.iges.or.jp/public/2006gl/index.html>.

Table 1-4: Summary of Key Categories for the United States (1990 and 2022) by Sector

CRT Code and Source/Sink Category	Greenhouse Gas	Approach 1 ^a				Approach 2 (includes uncertainty) ^a				2022 Emissions (MMT CO ₂ Eq.)
		Level Without LULUCF	Trend Without LULUCF	Level With LULUCF	Trend With LULUCF	Level Without LULUCF	Trend Without LULUCF	Level With LULUCF	Trend With LULUCF	
Energy										
1.A.3.b Transportation: Road	CO ₂	⊙	•	⊙	•	⊙	•	⊙	•	1,438.1
1.A.1 Stationary Combustion - Coal - Electricity Generation	CO ₂	⊙	•	⊙	•	⊙	•	⊙	•	851.5
1.A.1 Stationary Combustion - Natural Gas - Electricity Generation	CO ₂	⊙	•	⊙	•	⊙	•		•	659.3
1.A.2 Stationary Combustion - Natural Gas - Industrial	CO ₂	⊙	•	⊙	•	⊙	•	⊙	•	510.4
1.A.4.b Stationary Combustion - Natural Gas - Residential	CO ₂	⊙	•	⊙	•	⊙	•	⊙		272.0
1.A.2 Stationary Combustion - Oil - Industrial	CO ₂	⊙	•	⊙	•	⊙	•	⊙	•	247.6
1.A.4.a Stationary Combustion - Natural Gas - Commercial	CO ₂	⊙	•	⊙	•	⊙	•	•	•	192.3
1.A.3.a Transportation: Aviation	CO ₂	⊙	•	⊙	•	⊙		⊙		165.6
1.A.5 Non-Energy Use of Fuels	CO ₂	⊙		⊙		⊙	•	⊙		102.8
1.A.3.e Transportation: Other	CO ₂	⊙	•	⊙	•		•			69.3
1.A.4.a Stationary Combustion - Oil - Commercial	CO ₂	⊙	•	⊙	•					65.1
1.A.4.b Stationary Combustion - Oil - Residential	CO ₂	⊙	•	⊙	•					62.1

CRT Code and Source/Sink Category	Greenhouse Gas	Approach 1 ^a				Approach 2 (includes uncertainty) ^a				2022 Emissions (MMT CO ₂ Eq.)
		Level Without LULUCF	Trend Without LULUCF	Level With LULUCF	Trend With LULUCF	Level Without LULUCF	Trend Without LULUCF	Level With LULUCF	Trend With LULUCF	
1.A.2 Stationary Combustion - Coal - Industrial	CO ₂	⊙	•	⊙	•	⊙	•	○	•	43.0
1.A.3.d Transportation: Domestic Navigation	CO ₂	⊙		⊙						40.9
1.B.2 Natural Gas Systems	CO ₂	⊙		⊙						36.5
1.A.3.c Transportation: Railways	CO ₂	⊙		⊙						32.5
1.B.2 Petroleum Systems	CO ₂	•	•	•	•		•		•	22.0
1.A.1 Stationary Combustion - Oil - Electricity Generation	CO ₂	⊙	•	⊙	•	○	•		•	20.5
1.A.5 Stationary Combustion - Oil - U.S. Territories	CO ₂	○		○						17.0
1.A.5.b Transportation: Military	CO ₂		•		•					4.8
1.A.4.a Stationary Combustion - Coal - Commercial	CO ₂		•		•					1.4
1.A.4.b Stationary Combustion - Coal - Residential	CO ₂						•			NO
1.B.2 Natural Gas Systems	CH ₄	⊙	•	⊙	•	⊙	•	⊙	•	173.1
1.B.1 Fugitive Emissions from Coal Mining	CH ₄	⊙	•	⊙	•	⊙	•	⊙	•	43.6
1.B.2 Petroleum Systems	CH ₄	⊙	•	⊙	•	⊙	•	⊙	•	39.6
1.B.2 Abandoned Oil and Natural Gas Wells	CH ₄					⊙		⊙		8.5
1.A.4.b Stationary Combustion - Residential	CH ₄					⊙	•	⊙	•	4.3
1.A.1 Stationary Combustion - Coal - Electricity Generation	N ₂ O					⊙		•		18.2
1.A.3.b Transportation: Road	N ₂ O	○	•	○	•					8.9

CRT Code and Source/Sink Category	Greenhouse Gas	Approach 1 ^a				Approach 2 (includes uncertainty) ^a				2022 Emissions (MMT CO ₂ Eq.)
		Level Without LULUCF	Trend Without LULUCF	Level With LULUCF	Trend With LULUCF	Level Without LULUCF	Trend Without LULUCF	Level With LULUCF	Trend With LULUCF	
1.A.1 Stationary Combustion - Natural Gas - Electricity Generation	N ₂ O						•			3.4
Industrial Processes and Product Use										
2.A.1 Cement Production	CO ₂	⊙		⊙	•					41.9
2.C.1 Iron and Steel Production & Metallurgical Coke Production	CO ₂	⊙	•	⊙	•	⊙	•	○	•	40.7
2.B.8 Petrochemical Production	CO ₂	⊙	•	⊙	•					28.8
2.B.3 Adipic Acid Production	N ₂ O		•		•					2.1
2.F.1 Substitutes for Ozone Depleting Substances: Refrigeration and Air Conditioning	HFCs, PFCs	•	•	•	•	•	•	•	•	144.6
2.F.4 Substitutes for Ozone Depleting Substances: Aerosols	HFCs, PFCs		•		•	•	•	•	•	17.0
2.F.2 Substitutes for Ozone Depleting Substances: Foam Blowing Agents	HFCs, PFCs		•		•					11.7
2.B.9 Fluorochemical Production	PFCs, HFCs, SF ₆ , NF ₃	○	•	○	•	○	•	○	•	7.8
2.G Electrical Equipment	PFCs, SF ₆	○	•	○	•		•		•	5.1
2.C.3 Aluminum Production	PFCs	○	•	○	•					0.8
Agriculture										
3.A.1 Enteric Fermentation: Cattle	CH ₄	⊙	•	⊙	•	⊙	•	⊙	•	185.9
3.B.1 Manure Management: Cattle	CH ₄	•	•	•	•	•	•		•	37.7
3.B.4 Manure Management: Other Livestock	CH ₄	⊙		⊙		•				27.0
3.C Rice Cultivation	CH ₄	⊙		•		⊙		⊙		18.9

CRT Code and Source/Sink Category	Greenhouse Gas	Approach 1 ^a				Approach 2 (includes uncertainty) ^a				2022 Emissions (MMT CO ₂ Eq.)
		Level Without LULUCF	Trend Without LULUCF	Level With LULUCF	Trend With LULUCF	Level Without LULUCF	Trend Without LULUCF	Level With LULUCF	Trend With LULUCF	
3.D.1 Direct Emissions from Agricultural Soil Management	N ₂ O	⊙		⊙		⊙		⊙		262.5
3.D.2 Indirect Emissions from Applied Nitrogen	N ₂ O	⊙		⊙		⊙	•	⊙		28.3
Waste										
5.A Commercial Landfills	CH ₄	⊙	•	⊙	•	⊙	•	⊙	•	100.9
5.A Industrial Landfills	CH ₄	•		•		•	•			18.9
5.D Domestic Wastewater Treatment	CH ₄					○				13.6
5.D Domestic Wastewater Treatment	N ₂ O	•		•		⊙	•	⊙	•	21.4
Land Use, Land-Use Change, and Forestry										
4.E.2 Net Emissions from Land Converted to Settlements	CO ₂			⊙	•			⊙	•	68.2
4.B.2 Net Emissions from Land Converted to Cropland	CO ₂			⊙	•			⊙	•	35.1
4.C.2 Net Emissions from Land Converted to Grassland	CO ₂			⊙	•			⊙	•	25.6
4.C.1 Net Emissions from Grassland Remaining Grassland	CO ₂			○	•			⊙	•	13.4
4.B.1 Net Removals from Cropland Remaining Cropland	CO ₂			•	•			⊙	•	(31.7)
4.A.2 Net Removals from Land Converted to Forest Land	CO ₂			⊙				⊙		(100.3)
4.E.1 Net Removals from Settlements Remaining Settlements	CO ₂			⊙	•			⊙	•	(134.8)

CRT Code and Source/Sink Category	Greenhouse Gas	Approach 1 ^a				Approach 2 (includes uncertainty) ^a				2022 Emissions (MMT CO ₂ Eq.)
		Level Without LULUCF	Trend Without LULUCF	Level With LULUCF	Trend With LULUCF	Level Without LULUCF	Trend Without LULUCF	Level With LULUCF	Trend With LULUCF	
4.A.1 Net Removals from Forest Land Remaining Forest Land	CO ₂			⊙	•			⊙	•	(787.0)
4.D.1 Flooded Lands Remaining Flooded Lands	CH ₄			⊙						44.2
Subtotal of Key Categories Without LULUCF^b										6,169.7
Total Gross Emissions Without LULUCF										6,343.2
Percent of Gross Total Without LULUCF										97%
Subtotal of Key Categories With LULUCF^c										5,285.4
Total Net Emissions With LULUCF										5,488.9
Percent of Net Total With LULUCF										96%

NO (Not Occurring)

^a Symbols correspond to the year(s) in which a category is key: 1990 = ○; 2022 = •; 1990 and 2022 = ⊙.

^b Subtotal includes key categories from Level Approach 1 Without LULUCF, Level Approach 2 Without LULUCF, Trend Approach 1 Without LULUCF, and Trend Approach 2 Without LULUCF.

^c Subtotal includes key categories from Level Approach 1 With LULUCF, Level Approach 2 With LULUCF, Trend Approach 1 With LULUCF, and Trend Approach 2 With LULUCF.

Note: Parentheses indicate negative values (or sequestration).

1.6 Quality Assurance and Quality Control (QA/QC)

As part of efforts to achieve its stated goals for inventory quality, transparency, and credibility, the United States has developed a quality assurance and quality control plan designed to check, document, and improve the quality of its inventory over time. QA/QC activities on the *Inventory* are undertaken within the framework of the U.S. *Quality Assurance/Quality Control and Uncertainty Management Plan (QA/QC plan) for the U.S. Greenhouse Gas Inventory: Procedures Manual for QA/QC and Uncertainty Analysis*.

Key attributes of the QA/QC plan are summarized in Figure 1-2. These attributes include:

- *Procedures and Forms*: detailed and specific systems that serve to standardize the process of documenting and archiving QA/QC implementation and related information, as well as to guide the implementation of QA/QC and the analysis of uncertainty
- *Implementation of Procedures*: guidance on application of QA/QC procedures throughout the whole Inventory development process from initial data collection, through preparation of the emission and removal estimates, to publication of the *Inventory*, consistent with the *2006 IPCC Guidelines*.
- *Quality Assurance (QA)*: process for implementing expert and public reviews for both the inventory estimates and the Inventory report (which is the primary vehicle for disseminating the results of the inventory development process). The expert technical review conducted by the UNFCCC supplements these QA processes, consistent with the QA good practice and the *2006 IPCC Guidelines (IPCC 2006)*. See Section 1.3 for more details on these QA processes.
- *Quality Control (QC)*: application of *General (Tier 1) and Category-specific (Tier 2)* quality controls and checks, as recommended by *2006 IPCC Guidelines (IPCC 2006)*, along with consideration of secondary data and category-specific checks (additional Tier 2 QC) in parallel and coordination with the uncertainty assessment; the development of protocols and templates, which provides for more structured communication and integration with the suppliers of secondary information
- *General (Tier 1) and Category-specific (Tier 2) Checks*: quality controls and checks, as recommended by *IPCC Good Practice Guidance and 2006 IPCC Guidelines (IPCC 2006)*
- *Record Keeping*: provisions to track which procedures have been followed, the results of the QA/QC, uncertainty analysis, and feedback mechanisms for corrective action based on the results of the investigations which provide for continual data quality improvement and guided research efforts moving forward.
- *Multi-Year Implementation*: tracking the application of more involved QA/QC procedures which may take more than one cycle to fully implement, especially for category-specific QC, prioritizing key categories in conjunction with improvement planning (see Section 1.3).
- *Interaction and Coordination*: promoting communication within the EPA, across federal agencies and departments, state government programs, and research institutions and consulting firms involved in supplying data or preparing estimates for the *Inventory*. The QA/QC Management Plan itself is intended to be revised and reflect new information that becomes available as the program develops, methods are improved, or additional supporting documents become necessary.

Figure 1-2: Summary of Key QC Processes from U.S. QA/QC Plan

	Data Gathering	Data Documentation	Calculating Emissions	Cross-Cutting Coordination
Inventory Analyst	<ul style="list-style-type: none"> • Obtain data in electronic format (if possible) • Review data input/calculation workbooks <ul style="list-style-type: none"> ○ Avoid hardwiring ○ Use data validation ○ Protect cells • Develop automatic checkers for: <ul style="list-style-type: none"> ○ Outliers, negative values, or missing data ○ Variable types match values ○ Time series consistency • Maintain tracking tab for status of gathering efforts 	<ul style="list-style-type: none"> • Contact reports for non-electronic communications • Provide cell references for primary data elements • Obtain copies of all data sources • List and location of any working/external data or input/calculation workbooks • Document assumptions • Complete QA/QC checklists • CRF and summary tab links 	<ul style="list-style-type: none"> • Clearly label parameters, units, and conversion factors • Review data input/calculation workbooks integrity <ul style="list-style-type: none"> ○ Equations ○ Units ○ Inputs and outputs • Develop automated checkers for: <ul style="list-style-type: none"> ○ Input ranges ○ Calculations ○ Emission aggregation ○ Trend and IEF checks 	<ul style="list-style-type: none"> • Common starting versions for each inventory year • Utilize unalterable summary and CRF tab for each source data input/calculation workbook for linking to a master summary workbook • Follow strict version control procedures • Document QA/QC procedures
QA/QC Analyst	<ul style="list-style-type: none"> • Check input data for transcription errors • Inspect automatic checkers • Identify data input/calculation workbooks modifications that could provide additional QA/QC checks 	<ul style="list-style-type: none"> • Check citations in data input/calculation workbooks and text for accuracy and style • Check reference docket for new citations • Review documentation for any data / methodology changes • Complete QA/QC checklists • CRF and summary tab links 	<ul style="list-style-type: none"> • Reproduce calculations • Review time series consistency • Review changes in data/consistency with IPCC methodology 	

Box 1-3: Examples of Verification Activities

Consistent with IPCC guidance for national greenhouse gas inventories, verification activities include comparisons with emission or removal estimates prepared by other bodies and comparisons with estimates derived from fully independent assessments, e.g., atmospheric concentration measurements. Verification activities provide information to improve inventories and are part of the overall QA/QC system.

Use of Lower Tier Methods. The Paris Agreement/UNFCCC reporting guidelines require countries to complete a "top-down" reference approach for estimating CO₂ emissions from fossil fuel combustion in addition to their "bottom-up" sectoral methodology for purposes of verification. This estimation method uses alternative methodologies and different data sources than those contained in that section of the Energy chapter. The reference approach estimates fossil fuel consumption by adjusting national aggregate fuel production data for imports, exports, and stock changes rather than relying on end-user consumption surveys (see Annex 4 of this report). The reference approach assumes that once carbon-based fuels are brought into a national economy, they are either saved in some way (e.g., stored in products, kept in fuel stocks, or left unoxidized in ash) or combusted, and therefore the carbon in them is oxidized and released into the atmosphere. Accounting for actual consumption of fuels at the sectoral or sub-national level is not required.

Use of Ambient Measurements Systems for Validation of Emission Inventories. In following the Paris Agreement and UNFCCC reporting requirements to develop and submit national greenhouse gas emission inventories, the emissions and sinks presented in this report are organized by source and sink categories and calculated using internationally accepted methods provided by the IPCC.⁶⁶ Several recent studies have estimated emissions at the national or regional level with estimated results that sometimes differ from EPA's estimate of emissions. EPA has engaged with researchers on how remote sensing, ambient measurement, and inverse modeling techniques for estimating greenhouse gas emissions could assist in improving the understanding of inventory estimates. In working with the research community to improve national greenhouse gas inventories, EPA follows guidance from the IPCC on the use of measurements and modeling to validate emission inventories.⁶⁷ An area of particular interest in EPA's outreach efforts is how ambient measurement data can be used to assess estimates or potentially be incorporated into the *Inventory* in a manner consistent with this *Inventory* report's transparency of its calculation methodologies, and the ability of inverse modeling to attribute emissions and removals from remote sensing to anthropogenic sources, as defined by the IPCC for this report, versus natural sources and sinks.

The *2019 Refinement to the IPCC 2006 Guidelines for National Greenhouse Gas Inventories* (IPCC 2019) Volume 1 General Guidance and Reporting, Chapter 6: Quality Assurance, Quality Control and Verification notes that emission estimates derived from atmospheric concentration measurements can provide independent data sets as a basis for comparison with inventory estimates. The *2019 Refinement* provides guidance on conducting such comparisons (as summarized in Table 6.2 of IPCC [2019] Volume 1, Chapter 6) and provides guidance on using such comparisons to identify areas of improvement in national inventories (as summarized in Box 6.5 of IPCC [2019] Volume 1, Chapter 6). Further, it identified fluorinated gases as particularly suitable for such comparisons due their limited natural sources, their generally long atmospheric lifetimes, and well-understood loss mechanisms, which makes it relatively more straightforward to model their emission fluxes from observed mass quantities. Unlike emissions of CO₂, CH₄, and N₂O, emissions of fluorinated greenhouse gases are almost exclusively anthropogenic, meaning that the fluorinated greenhouse gas emission sources included in this *Inventory* account for the majority of the total U.S. emissions of these gases detectable in the atmosphere. This evaluation approach is also useful for gases and sources with larger uncertainties in available bottom-up inventory methods and data, such as emissions of CH₄, which are primarily from uncertain biological (e.g., enteric fermentation) and fugitive (e.g., natural gas production) activities.

In this *Inventory*, EPA includes the results from current and previous comparisons between fluorinated gas

⁶⁶ See <http://www.ipcc-nggip.iges.or.jp/public/index.html>.

⁶⁷ See http://www.ipcc-nggip.iges.or.jp/meeting/pdffiles/1003_Uncertainty%20meeting_report.pdf.

emissions inferred from atmospheric measurements and fluorinated gas emissions estimated based on bottom-up measurements and modeling. These comparisons, performed for HFCs and SF₆ respectively, are described under the QA/QC and Verification discussions in Chapter 4, Sections 4.25 Substitution of Ozone Depleting Substances and 4.26 Electrical Equipment in the IPPU chapter of this report.

Consistent with the *2019 Refinement*, a key element to facilitate such comparisons is a spatially-explicit (or gridded) emissions inventory as an input to inverse modeling. To improve the ability to compare methane emissions from the national-level greenhouse gas inventory with observation-based emission estimates, a team of researchers from U.S. EPA, SRON Netherlands Institute for Space Research, Harvard University, and Lawrence Berkely National Laboratory developed a time series of annual anthropogenic methane emissions maps with 0.1° x 0.1° (~10km x 10km) spatial resolution and monthly temporal resolution for the contiguous United States.⁶⁸ The gridded methane inventory is designed to be consistent with the *U.S. EPA Inventory of U.S. Greenhouse Gas Emissions and Sinks* estimates, which presents national totals for different source types.⁶⁹ The development of this gridded inventory is consistent with the recommendations contained in two National Academies of Science reports examining greenhouse gas emissions data (National Research Council 2010; National Academies of Sciences, Engineering, and Medicine 2018).

Finally, in addition to use of atmospheric concentration measurement data for comparison with Inventory data, information from top-down studies is directly incorporated in the Natural Gas Systems calculations to quantify emissions from certain well blowout events.

In addition, based on the national QA/QC plan for the *Inventory*, some sector, subsector and category-specific QA/QC and verification checks have been applied. These checks follow the procedures outlined in the national QA/QC plan, tailoring the procedures to the specific documentation and data files associated with individual sources. For each greenhouse gas emissions source or sink category included in this *Inventory*, a minimum of general or Tier 1 QC analysis has been undertaken. Where QC activities for a particular category go beyond the minimum general checks and include category-specific checks (Tier 2) or include verification, further explanation is provided within the respective source or sink category text. Similarly, responses or updates based on comments from the expert, public and the international technical expert reviews (e.g., UNFCCC) are also addressed within the respective source or sink category sections in each sectoral chapter and Annex 8.

The quality control activities described in the U.S. QA/QC plan occur throughout the inventory process; QA/QC is not separate from, but is an integral part of, preparing the *Inventory*. Quality control—in the form of both good practices (such as documentation procedures) and checks on whether good practices and procedures are being followed—is applied at every stage of inventory development and document preparation. In addition, quality assurance occurs during the expert review and the public review, in addition to the UNFCCC expert technical review. While all phases significantly contribute to improving inventory quality, the public review phase is also essential for promoting the openness of the inventory development process and the transparency of the inventory methods and underlying input data sources.

The QA/QC plan guides the process of ensuring inventory quality by describing data and methodology checks, developing processes governing peer review and public comments, and developing guidance on conducting an analysis of the uncertainty surrounding the emission and removal estimates. The QA/QC procedures also include feedback loops and provide for corrective actions that are designed to improve the inventory estimates over time.

⁶⁸ See <https://www.epa.gov/ghgemissions/us-gridded-methane-emissions>.

⁶⁹ See <https://www.epa.gov/ghgemissions/inventory-us-greenhouse-gas-emissions-and-sinks>.

1.7 Uncertainty Analysis

Emissions and removals calculated for the U.S. Inventory reflect best estimates for greenhouse gas source and sink categories in the United States and are continuously revised and improved as new information becomes available. Uncertainty assessment is an essential element of a complete and transparent emissions inventory because it helps inform and prioritize Inventory improvements. For the U.S. Inventory, uncertainty analyses are conducted for each source and sink category as well as for the uncertainties associated with the overall emission (current and base year) and trends estimates. These analyses reflect the quantitative uncertainty in the emission (and removal) estimates associated with uncertainties in their input parameters (e.g., activity data and EFs) and serve to evaluate the relative contribution of individual input parameter uncertainties to the overall Inventory, its trends, and each source and sink category.

The overall level and trend uncertainty estimates for total U.S. greenhouse gas emissions was developed using the IPCC Approach 2 uncertainty estimation methodology (assuming a Normal distribution for Approach 1 estimates), which employs a Monte Carlo stochastic simulation technique. The IPCC provides good practice guidance on two approaches—Approach 1 and Approach 2—to estimating uncertainty for both individual and combined source categories. Approach 2 quantifies uncertainties based on a distribution of emissions (or removals), built-up from repeated calculations of emission estimation models and the underlying input parameters, randomly selected according to their known distributions. Approach 2 methodology is applied to each individual source and sink category wherever data and resources are permitted and is also used to quantify the uncertainty in the overall Inventory and its Trends. Source and sink chapters in this report provide additional details on the uncertainty analysis conducted for each source and sink category. See Annex 7 of this report for further details on the U.S. process for estimating uncertainty associated with the overall emission (base and current year) and trends estimates. Consistent with IPCC (IPCC 2006), the United States has ongoing efforts to continue to improve the overall Inventory uncertainty estimates presented in this report.

The United States has also implemented many improvements over the last several years to reduce uncertainties across the source and sink categories and improve Inventory estimates. These improvements largely result from new data sources that provide more accurate data and/or increased data coverage, as well as methodological improvements. Following IPCC good practice, additional efforts to reduce Inventory uncertainties can occur through efforts to incorporate excluded emission and sink categories (see Annex 5), improve estimation methods, and collect more detailed, measured, and representative data. Individual category chapters and Annex 7 both describe current ongoing and planned Inventory and uncertainty analysis improvements. Consistent with IPCC (2006), the United States has ongoing efforts to continue to improve the category-specific uncertainty estimates presented in this report, largely prioritized by considering improvements categories identified as significant by the Key Category Analysis.

Estimates of quantitative uncertainty for the total U.S. greenhouse gas emissions in 1990 (base year) and 2022 are shown below in Table 1-5 and Table 1-6, respectively. The overall uncertainty surrounding the Total Net Emissions is estimated to be -6 to +6 percent in 1990 and -5 to +6 percent in 2022. When the LULUCF sector is excluded from the analysis the uncertainty is estimated to be -3 to +4 percent in 1990 and -2 to +4 percent in 2022.

Table 1-5: Estimated Overall Inventory Quantitative Uncertainty for 1990 (MMT CO₂ Eq. and Percent)

Gas	1990	Uncertainty Range Relative to Greenhouse Gas				Mean ^b	Standard Deviation ^b
	Emission Estimate (MMT CO ₂ Eq.)	Estimate ^a		Estimate ^a			
		(MMT CO ₂ Eq.)	(%)	(MMT CO ₂ Eq.)	(%)	(MMT CO ₂ Eq.)	
		Lower Bound ^c	Upper Bound ^c	Lower Bound	Upper Bound		
CO ₂	5,131.6	5,008.2	5,348.2	-2%	4%	5,098.2	88.0
CH ₄ ^d	871.7	731.3	948.4	-16%	9%	701.5	56.3
N ₂ O ^d	408.2	349.7	513.0	-14%	26%	434.8	41.6
PFC, HFC, SF ₆ , and NF ₃ ^d	125.5	108.6	152.9	-13%	22%	207.3	11.6
Total Gross Emissions	6,536.9	6,354.3	6,792.8	-3%	4%	6,441.8	113.3
LULUCF Emissions ^e	57.9	55.2	61.9	-5%	7%	68.7	1.7
LULUCF Carbon Stock Change Flux ^f	(1,034.7)	(1,296.1)	(845.3)	25%	-18%	(957.3)	116.7
LULUCF Sector Net Total^g	(976.7)	(1,237.7)	(787.8)	27%	-19%	(888.6)	116.7
Net Emissions (Sources and Sinks)	5,560.2	5,247.0	5,882.2	-6%	6%	5,553.3	161.4

- ^a The lower and upper bounds for emission estimates correspond to a 95 percent confidence interval, with the lower bound corresponding to 2.5th percentile and the upper bound corresponding to 97.5th percentile.
- ^b Mean value indicates the arithmetic average of the simulated emission estimates; standard deviation indicates the extent of deviation of the simulated values from the mean.
- ^c The lower and upper bound emission estimates for the sub-source categories do not sum to total emissions because the low and high estimates for total emissions were calculated separately through simulations.
- ^d The overall uncertainty estimates did not take into account the uncertainty in the GWP values for CH₄, N₂O and high GWP gases used in the Inventory emission calculations for 1990.
- ^e LULUCF emissions include the CH₄ and N₂O emissions reported for peatlands remaining peatlands, forest fires, drained organic soils, grassland fires, and coastal wetlands remaining coastal wetlands; CH₄ emissions from land converted to coastal wetlands, land converted to flooded land, and flooded land remaining flooded land; and N₂O emissions from forest soils and settlement soils.
- ^f LULUCF carbon stock change is the net C stock change from the following categories: forest land remaining forest land, land converted to forest land, cropland remaining cropland, land converted to cropland, grassland remaining grassland, land converted to grassland, wetlands remaining wetlands, land converted to wetlands, settlements remaining settlements, and land converted to settlements. Since the resulting flux is negative the signs of the resulting lower and upper bounds are reversed.
- ^g The LULUCF sector net total is the net sum of all CH₄ and N₂O emissions to the atmosphere plus net carbon stock changes.
- Notes: Total emissions (excluding emissions for which uncertainty was not quantified) are presented without LULUCF. Net emissions are presented with LULUCF. Totals may not sum due to independent rounding. Parentheses indicate net sequestration.

Table 1-6: Estimated Overall Inventory Quantitative Uncertainty for 2022 (MMT CO₂ Eq. and Percent)

Gas	2022 Emission Estimate	Uncertainty Range Relative to Greenhouse Gas				Mean ^b	Standard Deviation ^b
	(MMT CO ₂ Eq.)	Estimate ^a		Estimate ^a			
		(MMT CO ₂ Eq.)	(%)	(MMT CO ₂ Eq.)	(%)	(MMT CO ₂ Eq.)	
		Lower Bound ^c	Upper Bound ^c	Lower Bound	Upper Bound		
CO ₂	5,053.0	4,937.3	5,257.7	-2%	4%	5,095.2	81.9
CH ₄ ^d	702.4	604.3	803.1	-14%	14%	703.8	52.0
N ₂ O ^d	389.7	324.6	490.2	-17%	26%	399.5	42.3
PFC, HFC, SF ₆ , and NF ₃ ^d	198.1	182.8	217.5	-8%	10%	199.5	9.0

Total Gross Emissions	6,343.2	6,190.3	6,604.8	-2%	4%	6,397.9	106.3
LULUCF Emissions ^e	67.5	64.3	73.2	-5%	8%	68.6	2.3
LULUCF Carbon Stock Change Flux ^f	(921.8)	(1,158.6)	(748.7)	26%	-19%	(957.5)	105.3
LULUCF Sector Net Total^g	(854.3)	(1,090.3)	(680.5)	28%	-20%	(888.8)	105.3
Net Emissions (Sources and Sinks)	5,488.9	5,216.2	5,801.9	-5%	6%	5,509.0	150.6

^a The lower and upper bounds for emission estimates correspond to a 95 percent confidence interval, with the lower bound corresponding to 2.5th percentile and the upper bound corresponding to 97.5th percentile.

^b Mean value indicates the arithmetic average of the simulated emission estimates; standard deviation indicates the extent of deviation of the simulated values from the mean.

^c The lower and upper bound emission estimates for the sub-source categories do not sum to total emissions because the low and high estimates for total emissions were calculated separately through simulations.

^d The overall uncertainty estimates did not take into account the uncertainty in the GWP values for CH₄, N₂O and high GWP gases used in the Inventory emission calculations for 2022.

^e LULUCF emissions include the CH₄ and N₂O emissions reported for peatlands remaining peatlands, forest fires, drained organic soils, grassland fires, and coastal wetlands remaining coastal wetlands; CH₄ emissions from land converted to coastal wetlands, land converted to flooded land, and flooded land remaining flooded land; and N₂O emissions from forest soils and settlement soils.

^f LULUCF carbon stock change is the net C stock change from the following categories: forest land remaining forest land, land converted to forest land, cropland remaining cropland, land converted to cropland, grassland remaining grassland, land converted to grassland, wetlands remaining wetlands, land converted to wetlands, settlements remaining settlements, and land converted to settlements. Since the resulting flux is negative the signs of the resulting lower and upper bounds are reversed.

^g The LULUCF sector net total is the net sum of all CH₄ and N₂O emissions to the atmosphere plus net carbon stock changes.

Notes: Total emissions (excluding emissions for which uncertainty was not quantified) are presented without LULUCF. Net emissions are presented with LULUCF. Totals may not sum due to independent rounding. Parentheses indicate net sequestration.

The uncertainty for 2022 is similar to the uncertainty for 1990, though slightly lower. There have been some improvements in significant categories which do not necessarily reduce uncertainties as also acknowledged in the IPCC 2006 IPCC GL, p. 3.13 (e.g., improvements to estimates for Agricultural Soil Management over time, implications of methodological choice). For example, the 95 percent uncertainty bounds for nitrous oxide emissions from Agricultural Soil Management were increased from –25 percent to 29 percent in 1990 to –30 percent to 72 percent in 2022. Methodological and data quality improvements were also made for HFCs, PFCs, SF₆ and NF₃ this year but the uncertainties for these sources also slightly increased, better representing the limitations of existing emissions estimates. The methods and data for fossil fuel combustion categories, the most significant source, has not changed significantly and neither have uncertainties. It is also worth noting that some of the improvements to shift to use of GHGRP have been in less significant categories within the inventory (e.g., for IPPU). So, the overall uncertainty for latest year reflects these offsetting effects and trends within the uncertainty assessment.

In addition to the estimates of uncertainty associated with the current and base year estimates, Table 1-7 presents the estimates of inventory trend uncertainty. The 2006 IPCC Guidelines defines trend as the difference in emissions between the base year (i.e., 1990) and the current year (i.e., 2022) Inventory estimates. However, for purposes of understanding the concept of trend uncertainty, the trend is defined in this Inventory as the percentage change in the gross emissions (or net emissions) estimated for the current year, relative to the gross emission (or net emissions) estimated for the base year. The uncertainty associated with this trend is referred to as trend uncertainty and is reported as between -8 and 8 percent at the 95 percent confidence level between 1990 and 2022. This indicates a range of approximately -8 percent below and 8 percent above the trend estimate of -1.3 percent. See Annex 7 for trend uncertainty estimates for individual source and sink categories by gas.

Table 1-7: Quantitative Assessment of Trend Uncertainty (MMT CO₂ Eq. and Percent)

Gas	Base Year	2022	Emissions Trend	Trend Range ^b	
	Emissions ^a	Emissions		Trend Range ^b	
	(MMT CO ₂ Eq.)		(%)	(%)	
				Lower Bound	Upper Bound
CO ₂	5,131.6	5,053.0	-2%	-6%	3%
CH ₄	871.7	702.4	-19%	-32%	2%
N ₂ O	408.2	389.7	-5%	-30%	51%
HFCs, PFCs, SF ₆ , and NF ₃	125.5	198.1	58%	32%	95%
Total Gross Emissions^c	6,536.9	6,343.2	-3%	-7%	3%
LULUCF Emissions ^d	57.9	67.5	17%	6%	30%
LULUCF Carbon Stock Change Flux ^e	(1,034.7)	(921.8)	-11%	-35%	21%
LULUCF Sector Net Total^f	(976.7)	(854.3)	-13%	-37%	21%
Net Emissions (Sources and Sinks)^c	5,560.2	5,488.9	-1.3%	-8%	8%

^a Base Year is 1990 for all sources.

^b The trend range represents a 95 percent confidence interval for the emission trend, with the lower bound corresponding to 2.5th percentile value and the upper bound corresponding to 97.5th percentile value.

^c Totals exclude emissions for which uncertainty was not quantified.

^d LULUCF emissions include the CH₄ and N₂O emissions reported for peatlands remaining peatlands, forest fires, drained organic soils, grassland fires, and coastal wetlands remaining coastal wetlands; CH₄ emissions from land converted to coastal wetlands, land converted to flooded land, and flooded land remaining flooded land; and N₂O emissions from forest soils and settlement soils.

^e LULUCF carbon stock change is the net C stock change from the following categories: forest land remaining forest land, land converted to forest land, cropland remaining cropland, land converted to cropland, grassland remaining grassland, land converted to grassland, wetlands remaining wetlands, land converted to wetlands, settlements remaining settlements, and land converted to settlements.

^f The LULUCF sector net total is the net sum of all CH₄ and N₂O emissions to the atmosphere plus net carbon stock changes.

Notes: Totals may not sum due to independent rounding. Parentheses indicate net sequestration. Total emissions (excluding emissions for which uncertainty was not quantified) are presented without LULUCF. Net emissions are presented with LULUCF.

1.8 Completeness

This report, along with its accompanying CRTs, serves as a thorough assessment of the anthropogenic sources and sinks of greenhouse gas emissions for the United States for the time series 1990 through 2022. This report is intended to be comprehensive and includes the vast majority of emissions and removals identified as anthropogenic, consistent with IPCC methods and the Paris Agreement and UNFCCC reporting guidelines. In general, sources or sink categories not accounted for in this *Inventory* are excluded because they are not occurring in the United States and its territories, or because data are unavailable to develop an estimate and/or the categories were determined to be insignificant⁷⁰ in terms of overall national emissions per the Paris Agreement and UNFCCC reporting guidelines.

The United States is continually working to improve upon the understanding of such sources and sinks currently not included and seeking to find the data required to estimate related emissions and removals, focusing on categories that are anticipated to be significant. See Chapter 9 on Improvements and Recalculations for more

⁷⁰ See paragraph 32 in the Annex to Decision18/CMA.1 of the Paris Agreement reporting guidelines on national inventories that state "...emissions from a category should only be considered insignificant if the likely level of emissions is below 0.05 per cent of the national total GHG emissions, excluding LULUCF, or 500 kilotonnes of carbon dioxide equivalent (kt CO₂ eq), whichever is lower. The total national aggregate of estimated emissions for all gases from categories considered insignificant shall remain below 0.1 per cent of the national total GHG emissions, excluding LULUCF. Parties should use approximated activity data and default IPCC emission factors to derive a likely level of emissions for the respective category."

information on completeness and improvements implemented this cycle. As such improvements are implemented, new emission and removal estimates are quantified and included in the *Inventory*, improving completeness of national estimates. For a list of sources and sink categories not included and more information on significance of these categories, see Annex 5 and the respective category sections in each sectoral chapter of this report.

1.9 Organization of Report

In accordance with the Paris Agreement and UNFCCC reporting guidelines, this *Inventory* is grouped into five sector-specific chapters consistent with the Paris Agreement Common Reporting Tables (CRT),⁷¹ listed below in Table 1-8. In addition, the U.S. *Inventory* submission includes chapters on Trends in Greenhouse Gas Emissions, Other information, and Recalculations and Improvements to be considered consistent with the suggested outline or national inventory documents submitted under the Paris Agreement and UNFCCC.

Table 1-8: CRT/IPCC Sector Descriptions

Chapter (CRT and UNFCCC/IPCC Sector)	Activities Included
Energy	Emissions of all greenhouse gases resulting from stationary and mobile energy activities including fuel combustion and fugitive fuel emissions, and non-energy use of fossil fuels.
Industrial Processes and Product Use	Emissions resulting from industrial processes and product use of greenhouse gases.
Agriculture	Emissions from agricultural activities except fuel combustion, which is addressed under Energy.
Land Use, Land-Use Change, and Forestry	Emissions and removals of CO ₂ , and emissions of CH ₄ , and N ₂ O from land use, land-use change, and forestry.
Waste	Emissions from waste management activities.

Within each chapter, emissions are identified by the anthropogenic activity that is the source or sink of the greenhouse gas emissions being estimated (e.g., coal mining). Overall, the following organizational structure is consistently applied throughout this report:

Chapter/CRT/IPCC Sector: Overview of emissions and trends for each CRT/IPCC defined sector.

CRT Source or Sink Category: Description of category pathway and emission/removal trends based on IPCC methodologies, consistent with the Paris Agreement and UNFCCC reporting guidelines.

Methodology and Time-Series Consistency: Description of analytical methods (e.g., from *2006 IPCC Guidelines*, or country-specific methods) employed to produce emission estimates and identification of data references, primarily for activity data and emission factors, and a discussion of time-series consistency.

Uncertainty: A discussion and quantification of the uncertainty in emission estimates.

QA/QC and Verification: A discussion on steps taken to QA/QC and verify the emission estimates, consistent with the U.S. QA/QC plan, and any key QC findings.

Recalculations Discussion: A discussion of any data or methodological changes that necessitate a recalculation of previous years' emission estimates, and the impact of the recalculation on the emission estimates, if applicable.

Planned Improvements: A discussion on any category-specific planned improvements, if applicable.

⁷¹ See paragraph 50 in the Annex to Decision 18/CMA.1

Special attention is given to CO₂ from fossil fuel combustion relative to other sources because of its share of emissions and its dominant influence on emission trends. For example, each energy consuming end-use sector (i.e., residential, commercial, industrial, and transportation), as well as the electricity generation sector, is described individually. Additional information for certain source categories and other topics is also provided in several Annexes listed in Table 1-9.

Table 1-9: List of Annexes

ANNEX 1	Key Category Analysis
ANNEX 2	Methodology and Data for Estimating CO ₂ Emissions from Fossil Fuel Combustion
2.1.	Methodology for Estimating Emissions of CO ₂ from Fossil Fuel Combustion
2.2.	Methodology for Estimating the Carbon Content of Fossil Fuels
2.3.	Methodology for Estimating Carbon Emitted from Non-Energy Uses of Fossil Fuels
ANNEX 3	Methodological Descriptions for Additional Source or Sink Categories
3.1.	Methodology for Estimating Emissions of CH ₄ , N ₂ O, and Indirect Greenhouse Gases from Stationary Combustion
3.2.	Methodology for Estimating Emissions of CH ₄ , N ₂ O, and Indirect Greenhouse Gases from Mobile Combustion and Methodology for and Supplemental Information on Transportation-Related Greenhouse Gas Emissions
3.3.	Methodology for Estimating Emissions from Commercial Aircraft Jet Fuel Consumption
3.4.	Methodology for Estimating CH ₄ Emissions from Coal Mining
3.5.	Methodology for Estimating CH ₄ and CO ₂ Emissions from Petroleum Systems
3.6.	Methodology for Estimating CH ₄ Emissions from Natural Gas Systems
3.7.	Methodology for Estimating CO ₂ and N ₂ O Emissions from Incineration of Waste
3.8.	Methodology for Estimating Emissions from International Bunker Fuels used by the U.S. Military
3.9.	Methodology for Estimating HFC and PFC Emissions from Substitution of Ozone Depleting Substances
3.10.	Methodology for Estimating CH ₄ Emissions from Enteric Fermentation
3.11.	Methodology for Estimating CH ₄ and N ₂ O Emissions from Manure Management
3.12.	Methodology for Estimating N ₂ O Emissions, CH ₄ Emissions and Soil Organic C Stock Changes from Agricultural Lands (Cropland and Grassland)
3.13.	Methodology for Estimating Net Carbon Stock Changes in Forest Land Remaining Forest Land and Land Converted to Forest Land
3.14.	Methodology for Estimating CH ₄ Emissions from Landfills
ANNEX 4	IPCC Reference Approach for Estimating CO ₂ Emissions from Fossil Fuel Combustion
ANNEX 5	Assessment of the Sources and Sinks of Greenhouse Gas Emissions Not Included
ANNEX 6	Additional Information
6.1.	Global Warming Potential Values
6.2.	Ozone Depleting Substance Emissions
6.3.	Greenhouse Gas Precursors: Mapping of NEI categories to the Inventory
6.4.	Constants, Units, and Conversions
6.5.	Chemical Formulas
ANNEX 7	Uncertainty
7.1.	Overview
7.2.	Methodology and Results
7.3.	Reducing Uncertainty
7.4.	Planned Improvements
7.5.	Additional Information on Uncertainty Analyses by Source
ANNEX 8	QA/QC Procedures
8.1.	Background
8.2.	Purpose
8.3.	Assessment Factors
8.4.	Responses During the Review Process
ANNEX 9	Use of Greenhouse Gas Reporting Program (GHGRP) in Inventory